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<p>(21) International Application Number: <b>PCT/US97/07625</b></p> <p>(22) International Filing Date: <b>7 May 1997 (07.05.97)</b></p> <p>(30) Priority Data: 08/646,538                    8 May 1996 (08.05.96)                    US</p> <p>(71) Applicant: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, represented by THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES [US/US]; Bethesda, MD 20892 (US).</p> <p>(72) Inventors: PAVLAKIS, George, N.; 9 Purdue Court, Rockville, MD 20850 (US). GAITANARIS, George, A.; 9 Augustine Court, Gaithersburg, MD 20879 (US). STAUBER, Roland, H.; 1820 Hammond Court, Frederick, MD 21702 (US). VOURNAKIS, John, N.; 6 Carriage Lane, Hanover, NH 03755 (US).</p> <p>(74) Agents: WEBER, Ellen, L. et al.; Townsend and Townsend and Crew LLP, 8th floor, Two Embarcadero Center, San Francisco, CA 94111 (US).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p><b>Published</b> <i>With international search report.</i></p>	
<p>(54) Title: <b>MUTANT <i>AEQUOREA VICTORIA</i> FLUORESCENT PROTEINS HAVING INCREASED CELLULAR FLUORESCENCE</b></p> <p>(57) Abstract</p> <p>The present invention is directed to mutants of the jellyfish <i>Aequorea victoria</i> green fluorescent protein (GFP) having at least 5 and preferably greater than 20 times the specific green fluorescence of the wild type protein. In other embodiments, the invention comprises mutant blue fluorescent proteins (BFPs) that emit an enhanced blue fluorescence. The invention also encompasses the expression of nucleic acids that encode a mutant GFP or BFP in a wide variety of engineered host cells, and the isolation of engineered proteins having increased fluorescent activity. The novel mutants of the present invention allow for a significantly more sensitive detection of fluorescence in engineered host cells than is possible with GFP or with its known mutants. Thus, the mutant fluorescent proteins provided herein can be used as sensitive reporter molecules to detect the cell and tissue-specific expression and subcellular compartmentalization of GFP or BFP mutants, or of chimeric proteins comprising GFP or BFP mutants fused to a regulatory sequence or to a second protein sequence.</p>			

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HAVING INCREASED CELLULAR FLUORESCENCE**

FIELD OF THE INVENTION

This invention generally relates to novel proteins  
10 and their production which are useful for detecting gene  
expression and for visualizing the subcellular targeting and  
distribution of selected proteins and peptides, among other  
things. The invention specifically relates to mutations in  
the gene coding for the jellyfish *Aequorea victoria* green  
15 fluorescent protein ("GFP"), which mutations encode mutant GFP  
proteins having either an enhanced green or a blue  
fluorescence, and uses for them.

20                   **BACKGROUND OF THE INVENTION**

Green fluorescent protein ("GFP") is a monomeric  
protein of about 27 kDa which can be isolated from the  
bioluminescent jellyfish *Aequorea victoria*. When wild type  
GFP is illuminated by blue or ultraviolet light, it emits a  
25 brilliant green fluorescence. Similar to fluorescein  
isothiocyanate, GFP absorbs ultraviolet and blue light with a  
maximum absorbance at 395 nm and a minor peak of absorbance at  
470 nm, and emits green light with a maximum emission at 509  
nm with a minor peak at 540 nm. GFP fluorescence persists  
30 even after fixation with formaldehyde, and it is more stable  
to photobleaching than fluorescein.

The gene for GFP has been isolated and sequenced.  
Prasher, D. C. et al. (1992), "Primary structure of the  
*Aequorea victoria* green fluorescent protein," *Gene* 111:229-  
35 233. Expression vectors that comprise the GFP gene or cDNA  
have been introduced into a variety of host cells. These host  
cells include: Chinese hamster ovary (CHO) cells, human  
embryonic kidney cells (HEK293), COS-1 monkey cells, myeloma

cells, NIH 3T3 mouse fibroblasts, PtK1 cells, BHK cells, PC12 cells, Xenopus, leech, transgenic zebra fish, transgenic mice, Drosophila and several plants. The GFP molecules expressed by these different cells have a similar fluorescence as the native molecules, demonstrating that the GFP fluorescence does not require any species-specific cofactors or substrates. See, e.g., Baulcombe, D. et al. (1995), "Jellyfish green fluorescent protein as a reporter for virus infections," *The Plant Journal* 7:1045-1053; Chalfie, M. et al. (1994), "Green fluorescent protein as a marker for gene expression," *Science* 263:802-805; Inouye, S. & Tsuji, F. (1994), "Aequorea green fluorescent protein: expression of the gene and fluorescent characteristics of the recombinant protein," *FEBS Letters* 341:277-280; Inouye, S. & Tsuji, F. (1994), "Evidence for redox forms of the Aequorea green fluorescent protein," *FEBS Letters* 351:211-214; Kain, S. et al. (1995), "The green fluorescent protein as a reporter of gene expression and protein localization," *BioTechniques* (in press); Kitts, P. et al. (1995), "Green Fluorescent Protein (GFP): A novel reporter for monitoring gene expression in living organisms," *CLONTECHniques* X(1):1-3; Lo, D. et al. (1994), "Neuronal transfection in brain slices using particle-mediated gene transfer," *Neuron* 13:1263-1268; Moss, J. B. & Rosenthal, N. (1994), "Analysis of gene expression patterns in the embryonic mouse myotome with the green fluorescent protein, a new vital marker," *J. Cell. Biochem.*, Supplement 18D W161; Niedz, R. et al. (1995), "Green fluorescent protein: an in vivo reporter of plant gene expression," *Plant Cell Reports* 14:403-406; Wu, G.-I. et al. (1995), "Infection of frog neurons with vaccinia virus permits in vivo expression of foreign proteins," *Neuron* 14:681-684; Yu, J. & van den Engh, G. (1995), "Flow-sort and growth of single bacterial cells transformed with cosmid and plasmid vectors that include the gene for green-fluorescent protein as a visible marker," Abstracts of papers presented at the 1995 meeting on "Genome Mapping and Sequencing," Cold Spring Harbor, p. 293.

The active GFP chromophore is a hexapeptide which contains a cyclized Ser-dehydroTyr-gly trimer at positions 65-

67. This chromophore is only fluorescent when embedded within the intact GFP protein. Chromophore formation occurs post-translationally; nascent GFP is not fluorescent. The chromophore is thought to be formed by a cyclization reaction and an oxidation step that requires molecular oxygen.

5 Proteins can be fused to the amino (N-) or carboxy (C-) terminus of GFP. Such fused proteins have been shown to retain the fluorescent properties of GFP and the functional properties of the fusion partner. Bian, J. et al. (1995),  
10 "Nuclear localization of HIV-1 matrix protein P17: The use of A. victoria GFP in protein tagging and tracing," *FASEB J.* 9:AI279; Flach, J. et al. (1994), "A yeast RNA-binding protein shuttles between the nucleus and the cytoplasm," *Mol. Cell. Biol.* 14:8399-8407; Marshall, J. et al. (1995), "The  
15 jellyfish green fluorescent protein: a new tool for studying ion channel expression and function," *Neuron* 14:211-215; Olmsted, J. et al. (1994), "Green Fluorescent Protein (GFP) chimeras as reporters for MAP4 behavior in living cells," *Mol. Biol. of the Cell* 5:167a; Rizzuto, R. et al. (1995), "Chimeric  
20 green fluorescent protein as a tool for visualizing subcellular organelles in living cells," *Current Biol.* 5:635-642; Sengupta, P. et al. (1994), "The C. elegans gene odr-7 encodes an olfactory-specific member of the nuclear receptor superfamily," *Cell* 79:971-980; Stearns, T. (1995),  
25 "The green revolution," *Current Biol.* 5:262-264; Treinin, M. & Chalfie, M. (1995), "A mutated acetylcholine receptor subunit causes neuronal degeneration in C. elegans," *Neuron* 14:871-877; Wang, S. & Hazelrigg, T. (1994), "Implications for bcd mRNA localization from spatial distribution of exu protein in  
30 Drosophila oogenesis," *Nature* 369:400-403.

A number of GFP mutants have been reported.  
Delagrange, S. et al. (1995) "Red-shifted excitation mutants of the green fluorescent protein," *Bio/Technology* 13:151-154;  
Heim, R. et al. (1994) "Wavelength mutations and  
35 posttranslational autoxidation of green fluorescent protein," *Proc. Natl. Acad. Sci. USA* 91:12501-12504; Heim, R. et al. (1995), "Improved green fluorescence," *Nature* 373:663-664. Delgrave et al. (1995) *Bio/Technology* 13:151-154 isolated

mutants of cloned *Aequorea victoria* GFP that had red-shifted excitation spectra. Heim, R. et al. (1994) "Wavelength mutations and posttranslational autoxidation of green fluorescent protein," *Proc. Natl. Acad. Sci. USA* 91:12501-12504 reported a mutant (Tyr66 to His) having a blue fluorescence, which is herein designated BFP(Tyr<sub>67</sub>→His). These references have neither taught nor suggested that their mutations resulted in an increase in the cellular fluorescence of the mutant GFPs.

In general, the level of fluorescence of a protein expressed in a cell depends on several factors, such as number of copies made of the fluorescent protein, stability of the protein, efficiency of formation of the chromophore, and interactions with cellular solvents, solutes and structures. Although the fluorescent signal from wild type GFP or from the reported mutants is generally adequate for bulk detection of abundantly expressed GFP or of GFP-containing chimeras, it is inadequate for detecting transient low or constitutively low levels of expression, or for performing fine structural subcellular localizations. This limitation severely restricts the use of native GFP or of the reported mutants as a biochemical and structural marker for gene expression and morphological studies.

25

#### SUMMARY OF THE INVENTION

It is an object of the invention to provide engineered GFP-encoding nucleic acid sequences that encode modified GFP molecules having a greater cellular fluorescence than wild type GFP or prior described recombinant GFP.

It is a further object of this invention to provide recombinant vectors containing these modified GFP-encoding nucleic acid sequences, which vectors are capable of being inserted into a variety of cells (including mammalian and eukaryotic cells) and expressing the modified GFP.

It is also an object of this invention to provide host cells capable of providing useful quantities of homogeneous modified GFP.

It is yet another object of this invention to provide peptides that possess a greater cellular fluorescence than native GFP or unaltered recombinant GFP and that can be produced in large quantities in a laboratory, by a microorganism or by a cell in culture.

These and other objects of the invention have been accomplished by providing mutant GFP-encoding nucleic acids whose gene product exhibits an increased cellular fluorescence relative to naturally occurring or recombinantly produced wild type GFP ("wtGFP"). In some embodiments, the modified GFPs possess fluorescent activity that is 50-100 fold greater than that of unmodified GFP.

The modified proteins of the present invention are produced by making mutations in a genetic sequence that result in alterations in the amino acid sequence of the resulting gene product. Our starting material was a GFP-encoding nucleic acid wherein a codon encoding an additional nucleic acid was inserted at position 2 of the previously published GFP amino acid sequence (Chalfie et al., 1994), to introduce a useful restriction site. Due to the amino acid insertion at position 2 of the GFP amino acid sequence, our numbering of the GFP amino acids and description of the amino acid amutations is off by one as compared to the originally reported wild type GFP sequence (Prasher et al., 1992). Thus, amino acid 65 by our numbering corresponds to amino acid 64 of the originally reported wild type GFP, amino acid 168 corresponds to amino acid 167 of the originally reported wild type GFP, etc.

Using the modified wild type GFP described herein, a number of the unique mutants described herein derive from the discovery of an unplanned and unexpected mutation called "SG12", obtained in the course of site-directed mutagenesis experiments, wherein a phenylalanine at position 65 of wtGFP was converted to leucine. A mutant referred to as "SG11," which combined the phenylalanine 65 to leucine alteration with an isoleucine 168 to threonine substitution and a lysine 239 to asparagine substitution, gave a further enhanced fluorescence intensity. The lysine 239 to asparagine

substitution does not affect the fluorescence of GFP; indeed the C-terminal lysine or asparagine may be deleted without affecting fluorescence. A third and further improved GFP mutant was obtained by further mutating "SG11." This mutant 5 is referred to as "SG25" and, in addition to the SG11 mutations, contains an additional mutation, a substitution of a cysteine at position 66 for the serine normally found at that position in the sequence.

In addition, the invention encompasses novel GFP 10 mutants that emit a blue fluorescence. These blue mutants are derived from a mutation of the wild type GFP (Heim, R. et al. (1994) "Wavelength mutations and posttranslational autoxidation of green fluorescent protein," *Proc. Natl. Acad. Sci. USA* 91:12501-12504), in which histidine was substituted 15 for tyrosine at amino acid position 66. This mutant emits a blue fluorescence, i.e., it becomes a Blue Fluorescent Protein (BFP).

Novel BFP mutants having an enhanced blue 20 fluorescence were made by further modifying this BFP( $Tyr_{67}\rightarrow His$ ). The introduction of the same mutation used to generate SG12, (i.e., phenylalanine to leucine at position 65) into BFP( $Tyr_{67}\rightarrow His$ ) resulted in a new mutant having a brighter fluorescence, designated "SuperBlue-42" (SB42). A second independently generated mutation of BFP( $Tyr_{67}\rightarrow His$ ), in which a 25 valine at position 164 was converted to alanine, also emitted an enhanced blue fluorescent signal and is referred to as "SB49." A combination of the above two mutations resulted in "SB50", which exhibited an even greater fluorescence enhancement than either of the previous mutations.

The novel GFP and BFP mutants of this invention 30 allow for a significantly more sensitive detection of fluorescence in host cells than is possible with the wild type protein. Accordingly, the mutant GFPs provided herein can be used, among other things, as sensitive reporter molecules to detect the cell and tissue-specific expression and subcellular 35 compartmentalization of GFP or of chimeric proteins comprising GFP fused to a regulatory sequence or to a second protein sequence. In addition, these mutations make possible a

variety of one and two color protein assays to quantitate expression in mammalian cells.

5

#### DETAILED DESCRIPTION OF THE INVENTION

The present invention comprises mutant nucleic acids that encode engineered GFPs having a greater cellular fluorescence than either native GFP or unaltered ("wild type") recombinant GFP, and the mutant GFPs themselves. It further comprises a subset of mutant GFPs that are mutant blue fluorescent proteins ("BFPs") that are derived from a published BFP, designated BFP(Tyr<sub>67</sub>→His), wherein the mutant BFPs have a cellular fluorescence that is at least five times greater, preferably ten times greater, and most preferably 20 times greater than that of BFP(Tyr<sub>67</sub>→His). The invention also encompasses compositions such as vectors and cells that comprise either the mutant nucleic acids or the mutant protein gene products. The mutant GFP nucleic acids and proteins may be used to detect and quantify gene expression in living cells, and to detect and quantify tissue specific expression and subcellular distribution of GFP or of GFP fused to other proteins.

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##### I. General Definitions

30

Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Singleton et al. (1994) *Dictionary of Microbiology and Molecular Biology*, second edition, John Wiley and Sons (New York) provides one of skill with a general dictionary of many of the terms used in this invention. Although any methods and materials similar or equivalent to those described herein can be used in the practice or testing 35 of the present invention, the preferred methods and materials are described. For purposes of the present invention, the following terms are defined below.

The symbols, abbreviations and definitions used herein are set forth below:

	DNA, deoxyribonucleic acid
	RNA, ribonucleic acid
5	mRNA, messenger RNA
	cDNA, complementary DNA (enzymatically synthesized from an mRNA sequence)
	A-Adenine
	T-Thymine
10	G-Guanine
	C-Cytosine
	U-Uracil
	GFP, Green Fluorescent Protein
	BFP, Blue Fluorescent Protein
15	

Amino acids are sometimes referred to herein by the conventional one or three letter codes.

Wild type green fluorescent protein ("wtGFP") refers to the 239 amino acid sequence described by Chalfie et al., 20 Science 263, 802-805, 1994, the nucleotide sequence of which is set out as SEQ ID NO:1, and the amino acid sequence of which is set out as SEQ ID NO:2. This sequence differs from the original 238 amino acid GFP isolated from the bioluminescent jellyfish *Aequorea victoria* in that one amino acid has been inserted after position 2 of the 238 amino acid sequence. When reference in this application is made to an amino acid position of GFP, the position is made with reference to that described by Chalfie et al., *supra* and thus of SEQ ID NO:2.

30 The term "blue fluorescent protein" (BFP) refers to mutants of wtGFP wherein the tyrosine at position 67 is converted to a histidine, which mutants emit a blue fluorescence. The non-limiting prototype is herein designated BFP(Tyr<sub>67</sub>→His).

35 A shorthand designation for mutations that result in a change in amino acid sequence is the one or three letter code for the original amino acid, the number of the position of the amino acid in the wtGFP sequence, followed by the one or three letter code for the new amino acid. Thus, Phe65Leu or F65L both designate a mutation wherein the phenylalanine at position 65 of the wtGFP is converted to leucine.

40 Salts of any of the proteins described herein will naturally occur when such proteins are present in (or isolated

from) aqueous solutions of various pHs. All salts of peptides having the indicated biological activity are considered to be within the scope of the present invention. Examples include alkali, alkaline earth, and other metal salts of carboxylic acid residues, acid addition salts (e.g., HCl) of amino residues, and Zwitterions formed by reactions between carboxylic acid and amino acid residues within the same molecule.

The terms "bioluminescent" and "fluorescent" refer to the ability of GFP or of a derivative thereof to emit light ("emitted or fluorescent light") of a characteristic wavelength when excited by light which is generally of a characteristic and different wavelength than that used to generate the emission.

The term "cellular fluorescence" denotes the fluorescence of a GFP-derived protein of the present invention when expressed in a cell, especially a mammalian cell.

The term "nucleic acid" refers to a deoxyribonucleotide or ribonucleotide polymer in either single- or double-stranded form, and unless specifically limited, encompasses known analogues of natural nucleotides that hybridize to nucleic acids in a manner similar to naturally occurring nucleotides. Unless otherwise indicated, a particular nucleic acid sequence implicitly provides the complementary sequence thereof, as well as the sequence explicitly indicated. As used herein, the terms "nucleic acid" and "gene" are interchangeable, and they encompass the term "cDNA."

The phrase "a nucleic acid sequence encoding" refers to a nucleic acid which contains sequence information that, if translated, yields the primary amino acid sequence of a specific protein or peptide. This phrase specifically encompasses degenerate codons (i.e., different codons which encode a single amino acid) of the native sequence or sequences which may be introduced to conform with codon preference in a specific host cell.

The phrase "nucleic acid construct" denotes a nucleic acid that is composed of two or more nucleic acid

sequences that are derived from different sources and that are ligated together using methods known in the art.

The term "regulatory sequence" denotes all the non-coding elements of a nucleic acid sequence required for the correct and efficient expression of the "coding region" (i.e., the region that actually encodes the amino acid sequence of a peptide or protein), e.g., binding sites for polymerases and transcription factors, transcription and translation initiation and termination sequences, TATA box, a promoter to direct transcription, a ribosome binding site for translational initiation, polyadenylation sequences, enhancer elements.

The term "isolated" refers to material which is substantially or essentially free from components which normally accompany it as found in its native state (for example, a band on a gel). The isolated nucleic acids and the isolated proteins of this invention do not contain materials normally associated with their *in situ* environment, in particular, nuclear, cytosolic or membrane associated proteins or nucleic acids other than those nucleic acids which are indicated. The term "homogeneous" refers to a peptide or DNA sequence where the primary molecular structure (i.e., the sequence of amino acids or nucleotides) of substantially all molecules present in the composition under consideration is identical. The term "substantially" used in the preceding sentences preferably means at least 80% by weight, more preferably at least 95% by weight, and most preferably at least 99% by weight.

The nucleic acids of this invention, whether RNA, cDNA, genomic DNA, or a hybrid of the various combinations, are synthesized *in vitro* or are isolated from natural sources or recombinant clones. The nucleic acids claimed herein are present in transformed or transfected whole cells, in transformed or transfected cell lysates, or in a partially purified or substantially pure form. The nucleic acids of the present invention are obtained as homogeneous preparations. They may be prepared by standard techniques well known in the art, including selective precipitation with such substances as

ammonium sulfate, isopropyl alcohol, ethyl alcohol, and/or exclusion, ion exchange or affinity column chromatography, immunopurification methods, and others.

The phrase "conservatively modified variants thereof," when used with reference to a protein, denotes conservative amino acid substitutions in which both the original and the substituted amino acids have similar structure (e.g., the R group contains a carboxylic acid) and properties (e.g., the original and the substituted amino acids are acidic, such as glutamic and aspartic acid), such that the substitutions do not essentially alter specified properties of the protein, such as fluorescence. Amino acid substitutions that are conservative are well known in the art. The phrase "conservatively modified variants thereof," when used to describe a reference nucleic acid, denotes nucleic acids having nucleotide substitutions that yield degenerate codons for a given amino acid or that encode conservative amino acid substitutions, as compared to the reference nucleic acid.

The term "recombinant" or "engineered" when used with reference to a nucleic acid or a protein generally denotes that the composition or primary sequence of said nucleic acid or protein has been altered from the naturally occurring sequence using experimental manipulations well known to those skilled in the art. It may also denote that a nucleic acid or protein has been isolated and cloned into a vector, or that the nucleic acid that has been introduced into or expressed in a cell or cellular environment other than the cell or cellular environment in which said nucleic acid or protein may be found in nature. The phrase "engineered *Aequorea victoria* fluorescent protein" specifically encompasses a protein obtained by introducing one or more sequence alterations into the coding region of a nucleic acid that encodes wild type *Aequorea victoria* GFP, wherein the gene product of the engineered nucleic acid is a fluorescent protein recognized by antisera to wild type *Aequorea victoria* GFP.

The term "recombinant" or "engineered" when used with reference to a cell indicates that, as a result of

experimental manipulation, the cell replicates or expresses a nucleic acid or expresses a peptide or protein encoded by a nucleic acid, whose origin is exogenous to the cell.

Recombinant cells can express nucleic acids that are not found 5 within the native (non-recombinant) form of the cell.

Recombinant cells can also express nucleic acids found in the native form of the cell wherein the nucleic acids are re-introduced into the cell by artificial means.

The term "vector" denotes an engineered nucleic acid 10 construct that contains sequence elements that mediate the replication of the vector sequence and/or the expression of coding sequences present on the vector. Examples of vectors include eukaryotic and prokaryotic plasmids, viruses (for example, the HIV virus), cosmids, phagemids, and the like.

15 The term "operably linked" refers to functional linkage between a first nucleic acid (for example, an expression control sequence such as a promoter or an array of transcription factor binding sites) and a second nucleic acid sequence, wherein the expression control sequence directs 20 transcription of the nucleic acid corresponding to the second sequence. One or more selected isolated nucleic acids may be operably linked to a vector by methods known in the art.

"Transduction" or "transformation" denotes the process whereby exogenous extracellular DNA is introduced into 25 a cell, such that the cell is capable of replicating and or expressing the exogenous DNA. Generally, a selected nucleic acid is first inserted into a vector and the vector is then introduced into the cell. For example, plasmid DNA that is introduced under appropriate environmental conditions may 30 undergo replication in the transformed cell, and the replicated copies are distributed to progeny cells when cell division occurs. As a result, a new cell line is established, containing the plasmid and carrying the genetic determinants thereof. Transformation by a plasmid in this manner, where 35 the plasmid genes are maintained in the cell line by plasmid replication, occurs at high frequency when the transforming plasmid DNA is in closed loop form, and does not or rarely occurs if linear plasmid DNA is used.

All the patents and publications cited in this disclosure are indicative of the level of skill of those skilled in the art to which this invention pertains and are all herein individually incorporated by reference for all 5 purposes.

## II. The GFP Mutants and Their Expression

### A. The GFP mutants

10 The isolated nucleic acids reported here are those that encode an engineered protein derived from *Aequorea victoria* green fluorescent protein ("GFP") having a fluorescence at maximum emission that is at least five times greater, preferably ten times greater, and most preferably 15 twenty times greater than the fluorescence at maximum emission of wild type GFP. In one embodiment, a nucleic acid encodes for leucine at amino acid position 65. This amino acid position is important for the enhanced fluorescence. In another embodiment the engineered isolated GFP nucleic acid 20 also encodes for threonine at amino acid position 168. In an additional embodiment, the engineered isolated GFP nucleic acid further encodes for cysteine at amino acid position 66.

Also described here are GFP mutants that have enhanced blue fluorescent properties. These mutants have an 25 isolated nucleic acid that encode an engineered *Aequorea victoria* blue fluorescent protein that encodes for histidine at amino acid position 67, leucine at amino acid position 65 and has a cellular fluorescence that is at least five times greater, preferably 10 times greater, most preferably 20 times 30 greater than that of BFP(Tyr<sub>67</sub>->His). An alternative isolated BFP nucleic acid is one that encodes for an engineered *Aequorea victoria* blue fluorescent protein wherein the engineered BFP has histidine at amino acid position 67 and alanine at amino acid position 164. A third engineered 35 isolated BFP nucleic acid sequence is one that has histidine at amino acid position 67, leucine at amino acid position 65 and alanine at amino acid position 164.

The nucleic acid and amino acid sequences for the wild type GFP are set out in SEQ ID NO:1 and SEQ ID NO:2. The sequence is well-known, well-described and readily available for manipulation and use. Vectors bearing the nucleic acid sequence are commercially readily available from, for example, Clontech Laboratories, Inc., Clontech Laboratories, Inc., Palo Alto, CA. Clontech provides a line of reporter vectors for GFP, including the cDNA construct described by Chalfie, et al., *supra*, a promoterless GFP vector for monitoring the expression of cloned promoters in mammalian cells, and a series of vectors for creating fusion proteins to either the amino or carboxy terminus of GFP.

One of skill in the art will recognize many ways of generating alterations in a given nucleic acid sequence. Such well-known methods include site-directed mutagenesis, PCR amplification using degenerate oligonucleotides, exposure of cells containing the nucleic acid to mutagenic agents or radiation, chemical synthesis of a desired oligonucleotide (e.g., in conjunction with ligation and/or cloning to generate large nucleic acids) and other well-known techniques. See, e.g., Berger and Kimmel, *Guide to Molecular Cloning Techniques, Methods in Enzymology* Volume 152 Academic Press, Inc., San Diego, CA (Berger); Sambrook et al. (1989) *Molecular Cloning - A Laboratory Manual* (2nd ed.) Vol. 1-3, Cold Spring Harbor Laboratory, Cold Spring Harbor Press, NY, (Sambrook); and *Current Protocols in Molecular Biology*, F.M. Ausubel et al., eds., *Current Protocols*, a joint venture between Greene Publishing Associates, Inc. and John Wiley & Sons, Inc., (1994 Supplement) (Ausubel); Pirrung et al., U.S. Patent No. 5,143,854; and Fodor et al., *Science*, 251, 767-77 (1991). Product information from manufacturers of biological reagents and experimental equipment also provide information useful in known biological methods. Such manufacturers include the SIGMA Chemical Company (Saint Louis, MO), R&D systems (Minneapolis, MN), Pharmacia LKB Biotechnology (Piscataway, NJ), CLONTECH Laboratories, Inc. (Palo Alto, CA), Chem Genes Corp., Aldrich Chemical Company (Milwaukee, WI), Glen Research, Inc., GIBCO BRL Life Technologies, Inc.

(Gaithersberg, MD), Fluka Chemica-Biochemika Analytika (Fluka Chemie AG, Buchs, Switzerland), and Applied Biosystems (Foster City, CA), as well as many other commercial sources known to one of skill. Using these techniques, it is possible to 5 substitute at will any nucleotide in a nucleic acid that encodes any GFP or BFP disclosed herein or any amino acid in a GFP or BFP described herein for a predetermined nucleotide or amino acid. For example, it is possible to generate at will modified GFPs and BFP( $Tyr_{67} \rightarrow His$ )s that contain leucine at 10 position 65 and one or two or three additional mutations at any other position of the wtGFP or BFP( $Tyr_{67} \rightarrow His$ ).

The sequence of the cloned genes and synthetic oligonucleotides can be verified using the chemical degradation method of A.M. Maxam et al. (1980), *Methods in Enzymology* 65:499-560. The sequence can be confirmed after 15 the assembly of the oligonucleotide fragments into the double-stranded DNA sequence using the method of Maxam and Gilbert, *supra*, or the chain termination method for sequencing double-stranded templates of R.B. Wallace et al. (1981), *Gene*, 20 16:21-26. DNA sequencing may also be performed by the PCR-assisted fluorescent terminator method (ReadyReaction DyeDeoxy Terminator Cycle Sequencing Kit, ABI, Columbia, MD) according to the manufacturer's instructions, using the ABI 25 Model 373A DNA Sequencing System. Sequencing data is analyzed using the commercially available Sequencher program (Gene Codes, Gene Codes, Ann Arbor, MI).

**B. Expression of Mutant GFP**

Clearly, the nucleic acid sequences of the present 30 invention are excellent reporter sequences since the expressed proteins can be readily detected by fluorescence as described below. The sequences can be used in conjunction with any application appreciated to date for GFP and further in applications where a greater degree of fluorescence is 35 required. Expression of the sequences described herein whether expression is desired alone or in combination with other sequences of interest is described below.

Vectors to which selected foreign nucleic acids are operably linked may be used to introduce these selected nucleic acids into host cells and mediate their replication and/or expression. Cloning vectors are useful for replicating the foreign nucleic acids and obtaining clones of specific foreign nucleic acid-containing vectors. Expression vectors mediate the expression of the foreign nucleic acid. Some vectors are both cloning and expression vectors.

Once a nucleic acid is synthesized or isolated and inserted into a vector and cloned, one may express the nucleic acid in a variety of recombinantly engineered cells known to those of skill in the art. As used herein, "expression" refers to transcription of nucleic acids, either without or preferably with subsequent translation.

Expression of a mutant BFP or of wild type or mutant GFP can be enhanced by including multiple copies of the GFP-encoding nucleic acid in a transformed host, by selecting a vector known to reproduce in the host, thereby producing large quantities of protein from exogenous inserted DNA (such as pUC8, ptac12, or pIN-III-ompA1, 2, or 3), or by any other known means of enhancing peptide expression. In all cases, wtGFP or mutant GFPs will be expressed when the DNA sequence is functionally inserted into a vector. "Functionally inserted" means that it is inserted in proper reading frame and orientation. Typically, a GFP gene will be inserted downstream from a promoter and will be followed by a stop codon, although production as a hybrid protein followed by cleavage may be used, if desired.

Examples of cells which are suitable for the cloning and expression of the nucleic acids of the invention include bacteria, yeast, filamentous fungi, insect (especially employing baculoviral vectors), and mammalian cells, in particular cells capable of being maintained in tissue culture.

Host cells are competent or rendered competent for transformation by various means. There are several well-known methods of introducing DNA into animal cells. These include: calcium phosphate precipitation, fusion of the recipient cells

with bacterial protoplasts containing the DNA, treatment of the recipient cells with liposomes containing the DNA, DEAE dextran, receptor-mediated endocytosis, electroporation and micro-injection of the DNA directly into the cells.

It is expected that those of skill in the art are knowledgeable in the numerous systems available for cloning and expression of nucleic acids. In brief summary, the expression of natural or synthetic nucleic acids is typically achieved by operably linking a nucleic acid of interest to a promoter (which is either constitutive or inducible), and incorporating the construct into an expression vector. The vectors are suitable for replication and integration in prokaryotes, eukaryotes, or both. Typical cloning vectors contain transcription and translation terminators, transcription and translation initiation sequences, and promoters useful for regulation of the expression of the particular nucleic acid. The vectors optionally comprise generic expression cassettes containing at least one independent terminator sequence, sequences permitting replication of the cassette in eukaryotes, or prokaryotes, or both, (e.g., shuttle vectors) and selection markers for both prokaryotic and eukaryotic systems. See, e.g., Sambrook and Ausbel (both *supra*).

1. Expression in Prokaryotes

Prokaryotic systems for cloning and/or expressing engineered GFP or BFP proteins are available using *E. coli*, *Bacillus sp.* and *Salmonella* (Palva, I. et al. (1983), *Gene* 22:229-235; Mosbach, K. et al. (1983), *Nature* 302:543-545. To obtain high level expression in a prokaryotic system of a cloned nucleic acid such as those encoding engineered GFPs or BFPs, it is essential to construct expression vectors which contain, at a minimum, a strong promoter to direct transcription, a ribosome binding site for translational initiation, a transcription/translation terminator, a bacterial replicon, a nucleic acid encoding antibiotic resistance to permit selection of bacteria that harbor recombinant plasmids, and unique restriction sites in

nonessential regions of the plasmid to allow insertion of foreign nucleic acids. The particular antibiotic resistance gene chosen is not critical, any of the many resistance genes known in the art are suitable. Examples of regulatory regions 5 suitable for this purpose in *E. coli* are the promoter and operator region of the *E. coli* tryptophan biosynthetic pathway as described by Yanofsky, C. (1984), *J. Bacteriol.*, 158:1018-1024, and the leftward promoter of phage lambda ( $P_L$ ) as described by Herskowitz, I. and Hagen, D. (1980), *Ann. Rev. 10 Genet.*, 14:399-445 (1980).

The particular vector used to transport the genetic information into the cell is not particularly critical. Any of the conventional vectors used for replication, cloning and/or expression in prokaryotic cells may be used.

15 The foreign nucleic acid can be incorporated into a nonessential region of the host cell's chromosome. This is achieved by first inserting the nucleic acid into a vector such that it is flanked by regions of DNA homologous to the insertion site in the host chromosome. After introduction of 20 the vector into a host cell, the foreign nucleic acid is incorporated into the chromosome by homologous recombination between the flanking sequences and chromosomal DNA.

Detection of the expressed protein is achieved by methods known in the art as radioimmunoassays, or Western 25 blotting techniques or immunoprecipitation. Purification from *E. coli* can be achieved following procedures described in U.S. Patent No. 4,511,503.

## 2. Expression in Eukaryotes

30 Standard eukaryotic transfection methods are used to produce mammalian, yeast or insect cell lines which express large quantities of engineered GFP or BFP protein which are then purified using standard techniques. See, e.g., Colley et al. (1989), *J. Biol. Chem.* 264:17619-17622, and Guide to 35 Protein Purification, in Vol. 182 of *Methods in Enzymology* (Deutscher ed., 1990), D.A. Morrison (1977), *J. Bact.*, 132:349-351, or by J.E. Clark-Curtiss and R. Curtiss (1983).

*Methods in Enzymology* 101:347-362, Eds. R. Wu et al., Academic Press, New York.

The particular eukaryotic expression vector used to transport the genetic information into the cell is not particularly critical. Any of the conventional vectors used for expression in eukaryotic cells may be used. Expression vectors containing regulatory elements from eukaryotic viruses such as retroviruses are typically used. SV40 vectors include pSVT7 and pMT2. Vectors derived from bovine papilloma virus include pBV-1MTHA, and vectors derived from Epstein Barr virus include pHEBO, and p205. Other exemplary vectors include pMSG, pAV009/A<sup>+</sup>, pMT010/A<sup>+</sup>, pMAMneo-5, baculovirus pDSVE, and any other vector allowing expression of proteins under the direction of the SV-40 early promoter, SV-40 later promoter, metallothionein promoter, murine mammary tumor virus promoter, Rous sarcoma virus promoter, polyhedrin promoter, or other promoters shown effective for expression in eukaryotic cells.

The expression vector typically comprises a eukaryotic transcription unit or expression cassette that contains all the elements required for the expression of the engineered GFP or BFP DNA in eukaryotic cells. A typical expression cassette contains a promoter operably linked to the DNA sequence encoding a engineered GFP or BFP protein and signals required for efficient polyadenylation of the transcript.

Eukaryotic promoters typically contain two types of recognition sequences, the TATA box and upstream promoter elements. The TATA box, located 25-30 base pairs upstream of the transcription initiation site, is thought to be involved in directing RNA polymerase to begin RNA synthesis. The other upstream promoter elements determine the rate at which transcription is initiated.

Enhancer elements can stimulate transcription up to 1,000 fold from linked homologous or heterologous promoters. Enhancers are active when placed downstream or upstream from the transcription initiation site. Many enhancer elements derived from viruses have a broad host range and are active in a variety of tissues. For example, the SV40 early gene

enhancer is suitable for many cell types. Other enhancer/promoter combinations that are suitable for the present invention include those derived from polyoma virus, human or murine cytomegalovirus, the long term repeat from various retroviruses such as murine leukemia virus, murine or Rous sarcoma virus and HIV. See, *Enhancers and Eukaryotic Expression*, Cold Spring Harbor Press, Cold Spring Harbor, N.Y. 1983, which is incorporated herein by reference.

In the construction of the expression cassette, the promoter is preferably positioned about the same distance from the heterologous transcription start site as it is from the transcription start site in its natural setting. As is known in the art, however, some variation in this distance can be accommodated without loss of promoter function.

In addition to a promoter sequence, the expression cassette should also contain a transcription termination region downstream of the structural gene to provide for efficient termination. The termination region may be obtained from the same gene as the promoter sequence or may be obtained from different genes.

If the mRNA encoded by the structural gene is to be efficiently translated, polyadenylation sequences are also commonly added to the vector construct. Two distinct sequence elements are required for accurate and efficient polyadenylation: GU or U rich sequences located downstream from the polyadenylation site and a highly conserved sequence of six nucleotides, AAUAAA, located 11-30 nucleotides upstream. Termination and polyadenylation signals that are suitable for the present invention include those derived from SV40, or a partial genomic copy of a gene already resident on the expression vector.

In addition to the elements already described, the expression vector of the present invention may typically contain other specialized elements intended to increase the level of expression of cloned nucleic acids or to facilitate the identification of cells that carry the transfected DNA. For instance, a number of animal viruses contain DNA sequences that promote the extra chromosomal replication of the viral

genome in permissive cell types. Plasmids bearing these viral replicons are replicated episomally as long as the appropriate factors are provided by genes either carried on the plasmid or with the genome of the host cell.

5       The DNA sequence encoding the engineered GFP or BFP protein may typically be linked to a cleavable signal peptide sequence to promote secretion of the encoded protein by the transformed cell. Such signal peptides would include, among others, the signal peptides from tissue plasminogen activator,  
10 insulin, neuron growth factor, and juvenile hormone esterase of *Heliothis virescens*. Additional elements of the cassette may include enhancers and, if genomic DNA is used as the structural gene, introns with functional splice donor and acceptor sites.

15      The vector may or may not comprise a eukaryotic replicon. If a eukaryotic replicon is present, then the vector is amplifiable in eukaryotic cells using the appropriate selectable marker. If the vector does not comprise a eukaryotic replicon, no episomal amplification is  
20 possible. Instead, the transfected DNA integrates into the genome of the transfected cell, where the promoter directs expression of the desired nucleic acid.

25      The vectors usually comprise selectable markers which result in nucleic acid amplification such as the sodium, potassium ATPase, thymidine kinase, aminoglycoside phosphotransferase, hygromycin B phosphotransferase, xanthine-guanine phosphoribosyl transferase, CAD (carbamyl phosphate synthetase, aspartate transcarbamylase, and dihydroorotate), adenosine deaminase, dihydrofolate reductase,  
30 and asparagine synthetase and ouabain selection.

35      Alternatively, high yield expression systems not involving nucleic acid amplification are also suitable, such as using a baculovirus vector in insect cells, with the engineered GFP or BFP encoding sequence under the direction of the polyhedrin promoter or other strong baculovirus promoters.

The expression vectors of the present invention will typically contain both prokaryotic sequences that facilitate the cloning of the vector in bacteria as well as one or more

eukaryotic transcription units that are expressed only in eukaryotic cells, such as mammalian cells. The prokaryotic sequences are preferably chosen such that they do not interfere with the replication of the DNA in eukaryotic cells.

5 Any of the well known procedures for introducing foreign nucleotide sequences into host cells may be used. These include the use of calcium phosphate transfection, polybrene, protoplast fusion, electroporation, liposomes, microinjection, plasma vectors, viral vectors and any of the 10 other well known methods for introducing cloned genomic DNA, cDNA, synthetic DNA or other foreign nucleic acid material into a host cell (see Sambrook et al., *supra*). It is only necessary that the particular genetic engineering procedure utilized be capable of successfully introducing at least one 15 nucleic acid into the host cell which is capable of expressing the engineered GFP or BFP protein.

### **3. Expression in insect cells**

The baculovirus expression vector utilizes the 20 highly expressed and regulated *Autographa californica* nuclear polyhedrosis virus (AcMNPV) polyhedrin promoter modified for the insertion of foreign nucleic acids. Synthesis of polyhedrin protein results in the formation of occlusion bodies in the infected insect cell. The baculovirus vector 25 utilizes many of the protein modification, processing, and transport systems that occur in higher eukaryotic cells. The recombinant eukaryotic proteins expressed using this vector have been found in many cases to be, antigenically, 30 immunogenically, and functionally similar to their natural counterparts.

Briefly, a DNA sequence encoding an engineered GFP or BFP is inserted into a transfer plasmid vector in the proper orientation downstream from the polyhedrin promoter, and flanked on both ends with baculovirus sequences. Cultured 35 insect cells, commonly *Spodoptera frugiperda* cells, are transfected with a mixture of viral and plasmid DNAs. The virus that develop, some of which are recombinant virus that result from homologous recombination between the two DNAs, are

plated at 100-1000 plaques per plate. The plaques containing recombinant virus can be identified visually because of their ability to form occlusion bodies or by DNA hybridization. The recombinant virus is isolated by plaque purification. The 5 resulting recombinant virus, capable of expressing engineered GFP or BFP, is self-propagating in that no helper virus is required for maintenance or replication. After infecting an insect culture with recombinant virus, one can expect to find recombinant protein within 48-72 hours. The infection is 10 essentially lytic within 4-5 days.

There are a variety of transfer vectors into which the engineered GFP or BFP nucleic acid can be inserted. For a summary of transfer vectors see Luckow, V.A. and Summers, M.D. (1988), *Bio/Technology* 6:47-55. Preferred is the transfer 15 vector pAcUW21 described by Bishop, D.H.L. (1992) in *Seminars in Virology* 3:253-264.

#### 4. Retroviral Vectors

Retroviral vectors are particularly useful for 20 modifying eukaryotic cells because of the high efficiency with which the retroviral vectors transduce target cells and integrate into the target cell genome. Additionally, the retroviruses harboring the retroviral vector are capable of infecting cells from a wide variety of tissues.

Retroviral vectors are produced by genetically manipulating retroviruses. Retroviruses are RNA viruses because the viral genome is RNA. Upon infection, this genomic RNA is reverse transcribed into a DNA copy which is integrated into the chromosomal DNA of transduced cells with a high 25 degree of stability and efficiency. The integrated DNA copy is referred to as a provirus and is inherited by daughter cells as is any other gene. The wild type retroviral genome and the proviral DNA have three genes: the *gag*, the *pol* and the *env* genes, which are flanked by two long terminal repeat 30 (LTR) sequences. The *gag* gene encodes the internal structural (nucleocapsid) proteins; the *pol* gene encodes the RNA directed DNA polymerase (reverse transcriptase); and the *env* gene 35 encodes viral envelope glycoproteins. The 5' and 3' LTRs

serve to promote transcription and polyadenylation of virion RNAs. Adjacent to the 5' LTR are sequences necessary for reverse transcription of the genome (the tRNA primer binding site) and for efficient encapsulation of viral RNA into particles (the Psi site). See Mulligan, R.C. (1983), In: *Experimental Manipulation of Gene Expression*, M. Inouye (ed), 155-173; Mann, R. et al. (1983), *Cell*, 33:153-159; Cone, R.D. and R.C. Mulligan (1984), *Proceedings of the National Academy of Sciences, U.S.A.* 81:6349-6353.

The design of retroviral vectors is well known to one of skill in the art. See Singer, M. and Berg, P. *supra*. In brief, if the sequences necessary for encapsidation (or packaging of retroviral RNA into infectious virions) are missing from the viral genome, the result is a *cis* acting defect which prevents encapsidation of genomic RNA. However, the resulting mutant is still capable of directing the synthesis of all virion proteins. Retroviral genomes from which these sequences have been deleted, as well as cell lines containing the mutant genome stably integrated into the chromosome are well known in the art and are used to construct retroviral vectors. Preparation of retroviral vectors and their uses are described in many publications including European Patent Application EPA 0 178 220, U.S. Patent 4,405,712, Gilboa (1986), *Biotechniques* 4:504-512, Mann, et al. (1983), *Cell* 33:153-159, Cone and Mulligan (1984), *Proc. Natl. Acad. Sci. USA* 81:6349-6353, Eglitis, M.A., et al. (1988) *Biotechniques* 6:608-614, Miller, A.D. et al. (1989) *Biotechniques* 7:981-990, Miller, A.D. (1992) *Nature, supra*, Mulligan, R.C. (1993), *supra*. and Gould, B. et al., and International Patent Application No. WO 92/07943 entitled "Retroviral Vectors Useful in Gene Therapy." The teachings of these patents and publications are incorporated herein by reference.

The retroviral vector particles are prepared by recombinantly inserting the nucleic acid encoding engineered GFP or BFP into a retrovirus vector and packaging the vector with retroviral capsid proteins by use of a packaging cell line. The resultant retroviral vector particle is incapable

of replication in the host cell and is capable of integrating into the host cell genome as a proviral sequence containing the engineered GFP or BFP nucleic acid. As a result, the patient is capable of producing engineered GFP or BFP and 5 metabolize glycogen to completion.

Packaging cell lines are used to prepare the retroviral vector particles. A packaging cell line is a genetically constructed mammalian tissue culture cell line that produces the necessary viral structural proteins required 10 for packaging, but which is incapable of producing infectious virions. Retroviral vectors, on the other hand, lack the structural genes but have the nucleic acid sequences necessary for packaging. To prepare a packaging cell line, an infectious clone of a desired retrovirus, in which the 15 packaging site has been deleted, is constructed. Cells comprising this construct will express all structural proteins but the introduced DNA will be incapable of being packaged. Alternatively, packaging cell lines can be produced by transforming a cell line with one or more expression plasmids 20 encoding the appropriate core and envelope proteins. In these cells, the *gag*, *pol*, and *env* genes can be derived from the same or different retroviruses.

A number of packaging cell lines suitable for the present invention are available in the prior art. Examples of 25 these cell lines include Crip, GPE86, PA317 and PG13. See Miller et al. (1991), *J. Virol.* 65:2220-2224, which is incorporated herein by reference. Examples of other packaging cell lines are described in Cone, R. and Mulligan, R.C. (1984), *Proceedings of the National Academy of Sciences*, 30 U.S.A., 81:6349-6353 and in Danos, O. and R.C. Mulligan (1988), *Proceedings of the National Academy of Sciences*, U.S.A., 85:6460-6464, Eglitis, M.A., et al. (1988) *Biotechniques* 6:608-614, also all incorporated herein by reference.

Packaging cell lines capable of producing retroviral 35 vector particles with chimeric envelope proteins may be used. Alternatively, amphotropic or xenotropic envelope proteins,

such as those produced by PA317 and GPX packaging cell lines may be used to package the retroviral vectors.

Transforming cells with nucleic acids can involve, for example, incubating the cells with viral vectors (e.g., 5 retroviral or adeno-associated viral vectors) containing with cells within the host range of the vector. See, e.g., *Methods in Enzymology*, Vol. 185, Academic Press, Inc., San Diego, CA (D.V. Goeddel, ed.) (1990) or M. Krieger (1990), *Gene Transfer and Expression -- A Laboratory Manual*, Stockton Press, New 10 York, NY, and the references cited therein.

##### 5. Transformation with adeno-associated virus

Adeno associated viruses (AAVs) require helper viruses such as adenovirus or herpes virus to achieve 15 productive infection. In the absence of helper virus functions, AAV integrates (site-specifically) into a host cell's genome, but the integrated AAV genome has no pathogenic effect. The integration step allows the AAV genome to remain genetically intact until the host is exposed to the 20 appropriate environmental conditions (e.g., a lytic helper virus), whereupon it re-enters the lytic life-cycle. Samulski (1993), *Current Opinion in Genetic and Development* 3:74-80 and the references cited therein provides an overview of the AAV life cycle.

25 AAV-based vectors are used to transduce cells with target nucleic acids, e.g., in the *in vitro* production of nucleic acids and peptides, and in *in vivo* and *ex vivo* gene therapy procedures. See, West et al. (1987), *Virology* 160:38-47; Carter et al. (1989) U.S. Patent No. 4,797,368; Carter et 30 al. (1993), WO 93/24641; Kotin (1994), *Human Gene Therapy* 5:793-801; Muzychka (1994), *J. Clin. Invest.* 94:1351 and Samulski (*supra*) for an overview of AAV vectors.

Recombinant AAV vectors (rAAV vectors) deliver 35 foreign nucleic acids to a wide range of mammalian cells (Hermonat & Muzychka (1984), *Proc. Natl. Acad. Sci. USA* 81:6466-6470; Tratschin et al. (1985), *Mol. Cell Biol.* 5:3251-3260), integrate into the host chromosome (McLaughlin et al. (1988), *J. Virol.* 62:1963-1973), and show stable

expression of the transgene in cell and animal models (Flotte et al. (1993), *Proc. Natl. Acad. Sci. USA* 90:10613-10617). Moreover, unlike some retroviral vectors, rAAV vectors are able to infect non-dividing cells (Podsakoff et al. (1994), *J. Virol.* 68:5656-66; Flotte et al. (1994), *Am. J. Respir. Cell Mol. Biol.* 11:517-521). Further advantages of rAAV vectors include the lack of an intrinsic strong promoter, thus avoiding possible activation of downstream cellular sequences, and their naked eicosahedral capsid structure, which renders them stable and easy to concentrate by common laboratory techniques. rAAV vectors are used to inhibit, e.g., viral infection, by including anti-viral transcription cassettes in the rAAV vector which comprise an inhibitor of the invention.

15           **6. Expression in recombinant vaccinia virus-infected cells**

The nucleic acid encoding engineered GFP or BFP is inserted into a plasmid designed for producing recombinant vaccinia, such as pGS62, Langford, C.L. et al. (1986), *Mol. Cell. Biol.* 6:3191-3199. This plasmid consists of a cloning site for insertion of foreign nucleic acids, the P7.5 promoter of vaccinia to direct synthesis of the inserted nucleic acid, and the vaccinia TK gene flanking both ends of the foreign nucleic acid.

25           When the plasmid containing the engineered GFP or BFP nucleic acid is constructed, the nucleic acid can be transferred to vaccinia virus by homologous recombination in the infected cell. To achieve this, suitable recipient cells are transfected with the recombinant plasmid by standard calcium phosphate precipitation techniques into cells already infected with the desirable strain of vaccinia virus, such as Wyeth, Lister, WR or Copenhagen. Homologous recombination occurs between the TK gene in the virus and the flanking TK gene sequences in the plasmid. This results in a recombinant 30 virus with the foreign nucleic acid inserted into the viral TK gene, thus rendering the TK gene inactive. Cells containing recombinant viruses are selected by adding medium containing 35 S-bromodeoxyuridine, which is lethal for cells expressing a TK gene.

Confirmation of production of recombinant virus is achieved by DNA hybridization using cDNA encoding the engineered GFP or BFP and by immunodetection techniques using antibodies specific for the expressed protein. Virus stocks 5 may be prepared by infection of cells such as HeLa S3 spinner cells and harvesting of virus progeny.

#### 7. Expression in cell cultures

GFP- or BFP-encoding nucleic acids can be ligated to 10 various expression vectors for use in transforming host cell cultures. The culture of cells used in conjunction with the present invention is well known in the art. Freshney (1994) (*Culture of Animal Cells, a Manual of Basic Technique, third edition* Wiley-Liss, New York), Kuchler et al. (1977) 15 *Biochemical Methods in Cell Culture and Virology*, Kuchler, R.J., Dowden, Hutchinson and Ross, Inc., and the references cited therein provides a general guide to the culture of cells. Illustrative cell cultures useful for the production of recombinant proteins include cells of insect or mammalian 20 origin. Mammalian cell systems often will be in the form of monolayers of cells, although mammalian cell suspensions are also used. Illustrative examples of mammalian cell lines include monocytes, lymphocytes, macrophage, VERO and HeLa cells, Chinese hamster ovary (CHO) cell lines, WI38, BHK, 25 Cos-7 or MDCK cell lines (see, e.g., Freshney, supra).

Cells of mammalian origin are illustrative of cell cultures useful for the production of the engineered GFP or BFP . Mammalian cell systems often will be in the form of monolayers of cells although mammalian cell suspensions may 30 also be used. Illustrative examples of mammalian cell lines include VERO and HeLa cells, Chinese hamster ovary (CHO) cell lines, WI38, BHK, COS-7 or MDCK cell lines.

As indicated above, the vector, e.g., a plasmid, which is used to transform the host cell, preferably contains 35 DNA sequences to initiate transcription and sequences to control the translation of the engineered GFP or BFP nucleic acid sequence. These sequences are referred to as expression control sequences. Illustrative expression control sequences

are obtained from the SV-40 promoter (*Science* 222:524-527, (1983)), the CMV i.e. Promoter (*Proc. Natl. Acad. Sci.* 81:659-663, (1984)) or the metallothionein promoter (*Nature* 296:39-42, (1982)). The cloning vector containing the expression control sequences is cleaved using restriction enzymes and adjusted in size as necessary or desirable and ligated with sequences encoding the engineered GFP or BFP protein by means well known in the art.

The vectors for transforming cells in culture typically contain gene sequences to initiate transcription and translation of the engineered GFP or BFP gene. These sequences need to be compatible with the selected host cell. In addition, the vectors preferably contain a marker to provide a phenotypic trait for selection of transformed host cells such as dihydrofolate reductase or metallothionein. Additionally, a vector might contain a replicative origin.

As mentioned above, when higher animal host cells are employed, polyadenylation or transcription terminator sequences from known mammalian genes need to be incorporated into the vector. An example of a terminator sequence is the polyadenylation sequence from the bovine growth hormone gene. Sequences for accurate splicing of the transcript may also be included. An example of a splicing sequence is the VP1 intron from SV40 (Sprague, J. et al. (1983), *J. Virol.* 45: 773-781).

Additionally gene sequences to control replication in the host cell may be incorporated into the vector such as those found in bovine papilloma virus type-vectors. Saveria-Campo, M. (1985), "Bovine Papilloma virus DNA a Eukaryotic Cloning Vector" in DNA Cloning Vol.II a Practical Approach Ed. D.M. Glover, IRL Press, Arlington, Virginia pp. 213-238.

The transformed cells are cultured by means well known in the art. For example, as published in Kuchler, R.J. et al., (1977), *Biochemical Methods in Cell Culture and Virology*.

In addition to the above general procedures which can be used for preparing recombinant DNA molecules and transformed unicellular organisms in accordance with the

practices of this invention, other known techniques and modifications thereof can be used in carrying out the practice of the invention. Any known system for expression of isolated genes is suitable for use in the present invention. For example, viral expression systems such as the baculovirus expression system are specifically contemplated within the scope of the invention. Many recent U.S. patents disclose plasmids, genetically engineering microorganisms, and methods of conducting genetic engineering which can be used in the practice of the present invention. For example, U.S. Pat. No. 4,273,875 discloses a plasmid and a process of isolating the same. U.S. Pat. No. 4,304,863 discloses a process for producing bacteria by genetic engineering in which a hybrid plasmid is constructed and used to transform a bacterial host. U.S. Pat. No. 4,419,450 discloses a plasmid useful as a cloning vehicle in recombinant DNA work. U.S. Pat. No. 4,362,867 discloses recombinant cDNA construction methods and hybrid nucleotides produced thereby which are useful in cloning processes. U.S. Pat. No. 4,403,036 discloses genetic reagents for generating plasmids containing multiple copies of DNA segments. U.S. Pat. No. 4,363,877 discloses recombinant DNA transfer vectors. U.S. Pat. No. 4,356,270 discloses a recombinant DNA cloning vehicle and is a particularly useful disclosure for those with limited experience in the area of genetic engineering since it defines many of the terms used in genetic engineering and the basic processes used therein. U.S. Pat. No. 4,336,336 discloses a fused gene and a method of making the same. U.S. Pat. No. 4,319,629 discloses plasmid vectors and the production and use thereof. U.S. Pat. No. 4,332,901 discloses a cloning vector useful in recombinant DNA. Although some of these patents are directed to the production of a particular gene product that is not within the scope of the present invention, the procedures described therein can easily be modified to the practice of the invention described in this specification by those skilled in the art of genetic engineering. Transferring the isolated GFP cDNA to other expression vectors will produce constructs which

improve the expression of the GFP polypeptide in *E. coli* or express GFP in other hosts.

### **III. Detection of GFP and BFP Nucleic Acids and Proteins**

5

#### **A. General detection methods**

The nucleic acids and proteins of the invention are detected, confirmed and quantified by any of a number of means well known to those of skill in the art. The unique quality 10 of the inventive expressed proteins here is that they provide an enhanced fluorescence which can be readily and easily observed. Fluorescence assays for the expressed proteins are described in detail below. Other general methods for detecting both nucleic acids and corresponding proteins 15 include analytic biochemical methods such as spectrophotometry, radiography, electrophoresis, capillary electrophoresis, high performance liquid chromatography (HPLC), thin layer chromatography (TLC), hyperdiffusion chromatography, and the like, and various immunological 20 methods such as fluid or gel precipitin reactions, immunodiffusion (single or double), immunoelectrophoresis, radioimmunoassays (RIAs), enzyme-linked immunosorbent assays (ELISAs), immunofluorescent assays, and the like. The detection of nucleic acids proceeds by well known methods such 25 as Southern analysis, northern analysis, gel electrophoresis, PCR, radiolabeling, scintillation counting, and affinity chromatography.

A variety of methods of specific DNA and RNA measurement using nucleic acid hybridization techniques are 30 known to those of skill in the art. For example, one method for evaluating the presence or absence of engineered GFP or BFP DNA in a sample involves a Southern transfer. Southern *et al.* (1975), *J. Mol. Biol.* 98:503. Briefly, the digested genomic DNA is run on agarose slab gels in buffer and 35 transferred to membranes. Hybridization is carried out using the probes discussed above. Visualization of the hybridized portions allows the qualitative determination of the presence or absence of engineered GFP or BFP genes.

Similarly, a Northern transfer may be used for the detection of engineered GFP or BFP mRNA in samples of RNA from cells expressing the engineered GFP or BFP gene. In brief, the mRNA is isolated from a given cell sample using an acid guanidinium-phenol-chloroform extraction method. The mRNA is then electrophoresed to separate the mRNA species and the mRNA is transferred from the gel to a nitrocellulose membrane. As with the Southern blots, labeled probes are used to identify the presence or absence of the engineered GFP or BFP transcript.

The selection of a nucleic acid hybridization format is not critical. A variety of nucleic acid hybridization formats are known to those skilled in the art. For example, common formats include sandwich assays and competition or displacement assays. Hybridization techniques are generally described in "Nucleic Acid Hybridization, A Practical Approach," Ed. Hames, B.D. and Higgins, S.J., IRL Press, 1985; Gall and Pardue (1969), Proc. Natl. Acad. Sci. USA 63:378-383; and John, Burnstein and Jones (1969), Nature 223:582-587.

For example, sandwich assays are commercially useful hybridization assays for detecting or isolating nucleic acid sequences. Such assays utilize a "capture" nucleic acid covalently immobilized to a solid support and labelled "signal" nucleic acid in solution. The clinical sample will provide the target nucleic acid. The "capture" nucleic acid and "signal" nucleic acid probe hybridize with the target nucleic acid to form a "sandwich" hybridization complex. To be effective, the signal nucleic acid cannot hybridize with the capture nucleic acid.

The nucleic acid sequences used in this invention can be either positive or negative probes. Positive probes bind to their targets and the presence of duplex formation is evidence of the presence of the target. Negative probes fail to bind to the suspect target and the absence of duplex formation is evidence of the presence of the target. For example, the use of a wild type specific nucleic acid probe or PCR primers may act as a negative probe in an assay sample where only the mutant engineered GFP or BFP is present.

Labelled signal nucleic acids, whether those described herein or others known in the art are used to detect hybridization. Complementary nucleic acids or signal nucleic acids may be labelled by any one of several methods typically used to detect the presence of hybridized polynucleotides.

5 One common method of detection is the use of autoradiography with  $^3\text{H}$ ,  $^{125}\text{I}$ ,  $^{35}\text{S}$ ,  $^{14}\text{C}$ , or  $^{32}\text{P}$ -labelled probes or the like. Other labels include ligands which bind to labelled

10 antibodies, fluorophores, chemiluminescent agents, enzymes, and antibodies which can serve as specific binding pair members for a labelled ligand.

Detection of a hybridization complex may require the binding of a signal generating complex to a duplex of target and probe polynucleotides or nucleic acids. Typically, such binding occurs through ligand and anti-ligand interactions as between a ligand-conjugated probe and an anti-ligand conjugated with a signal. The binding of the signal generation complex is also readily amenable to accelerations by exposure to ultrasonic energy.

20 The label may also allow indirect detection of the hybridization complex. For example, where the label is a hapten or antigen, the sample can be detected by using antibodies. In these systems, a signal is generated by attaching fluorescent or enzyme molecules to the antibodies or

25 in some cases, by attachment to a radioactive label. (Tijssen, P. (1985), "Practice and Theory of Enzyme Immunoassays," *Laboratory Techniques in Biochemistry and Molecular Biology*, Burdon, R.H., van Knippenberg, P.H., Eds., Elsevier, pp. 9-20.)

30 The sensitivity of the hybridization assays may be enhanced through use of a nucleic acid amplification system which multiplies the target nucleic acid being detected. *In vitro* amplification techniques suitable for amplifying sequences for use as molecular probes or for generating

35 nucleic acid fragments for subsequent subcloning are known. Examples of techniques sufficient to direct persons of skill through such *in vitro* amplification methods, including the polymerase chain reaction (PCR) the ligase chain reaction

(LCR), Q $\beta$ -replicase amplification and other RNA polymerase mediated techniques (e.g., NASBA) are found in Berger, Sambrook, and Ausubel, as well as Mullis et al. (1987), U.S. Patent No. 4,683,202; *PCR Protocols A Guide to Methods and Applications* (Innis et al., eds) Academic Press Inc. San Diego, CA (1990) (Innis); Arnheim & Levinson (October 1, 1990), *Chem. Eng. News* 36-47; J. NIH Res. (1991) 3:81-94; (Kwoh et al. (1989), *Proc. Natl. Acad. Sci. USA* 86:1173; Guatelli et al. (1990), *Proc. Natl. Acad. Sci. USA* 87:1874; Lomell et al. (1989), *J. Clin. Chem.* 35:1826; Landegren et al. (1988), *Science* 241:1077-1080; Van Brunt (1990), *Biotechnology* 8:291-294; Wu and Wallace (1989), *Gene* 4:560; Barringer et al. (1990), *Gene* 89:117, and Sooknanan and Malek (1995), *Biotechnology* 13:563-564. Improved methods of cloning in vitro amplified nucleic acids are described in Wallace et al., U.S. Pat. No. 5,426,039. Other methods recently described in the art are the nucleic acid sequence based amplification (NASBA™, Cangene, Mississauga, Ontario) and Q Beta Replicase systems. These systems can be used to directly identify mutants where the PCR or LCR primers are designed to be extended or ligated only when a select sequence is present. Alternatively, the select sequences can be generally amplified using, for example, nonspecific PCR primers and the amplified target region later probed for a specific sequence indicative of a mutation.

Oligonucleotides for use as probes, e.g., in in vitro amplification methods, for use as gene probes, or as inhibitor components are typically synthesized chemically according to the solid phase phosphoramidite triester method described by Beaucage and Caruthers (1981), *Tetrahedron Letts.* 22(20):1859-1862, e.g., using an automated synthesizer, as described in Needham-VanDevanter et al. (1984), *Nucleic Acids Res.* 12:6159-6168. Purification of oligonucleotides, where necessary, is typically performed by either native acrylamide gel electrophoresis or by anion-exchange HPLC as described in Pearson and Regnier (1983), *J. Chrom.* 255:137-149. The sequence of the synthetic oligonucleotides can be verified using the chemical degradation method of Maxam and Gilbert

(1980) in Grossman and Moldave (eds.) Academic Press, New York, *Methods in Enzymology* 65:499-560.

An alternative means for determining the level of expression of the engineered GFP or BFP gene is *in situ* hybridization. *In situ* hybridization assays are well known and are generally described in Angerer et al. (1987), *Methods Enzymol.* 152:649-660. In an *in situ* hybridization assay cells are fixed to a solid support, typically a glass slide. If DNA is to be probed, the cells are denatured with heat or alkali. The cells are then contacted with a hybridization solution at a moderate temperature to permit annealing of engineered GFP or BFP specific probes that are labelled. The probes are preferably labelled with radioisotopes or fluorescent reporters.

15

#### B. Fluorescence Assay

When a fluorophore such as protein that is capable of fluorescing is exposed to a light of appropriate wavelength, it will absorb and store light and then release the stored light energy. The range of wavelengths that a fluorophore is capable of absorbing is the excitation spectrum and the range of wavelengths of light that a fluorophore is capable of emitting is the emission or fluorescence spectrum. The excitation and fluorescence spectra for a given fluorophore usually differ and may be readily measured using known instruments and methods. For example, scintillation counters and photometers (e.g. luminometers), photographic film, and solid state devices such as charge coupled devices, may be used to detect and measure the emission of light.

The nucleic acids, vectors, mutant proteins provided herein, in combination with well known techniques for over-expressing recombinant proteins, make it possible to obtain unlimited supplies of homogeneous mutant GFPs and BFPs. These modified GFPs or BFPs having increased fluorescent activity replace wtGTP or other currently employed tracers in existing diagnostic and assay systems. Such currently employed tracers include radioactive atoms or molecules and color-producing enzymes such as horseradish peroxidase.

The benefits of using the mutants of the present invention are at least four-fold: the modified GFPs and BFPs are safer than radioactive-based assays, modified GFPs and BFPs can be assayed quickly and easily, and large numbers of samples can be handled simultaneously, reducing overall handling and increasing efficiency. Of great significance, the expression and subcellular distribution of the fluorescent proteins within cells can be detected in living tissues without any other experimental manipulation than to placing the cells on a slide and viewing them through a fluorescence microscope. This represents a vast improvement over methods of immunodetection that require fixation and subsequent labelling.

The modified GFPs and BFPs of the present invention can be used in standard assays involving a fluorescent marker. For example, ligand-ligator binding pairs that can be modified with the mutants of the present invention without disrupting the ability of each to bind to the other can form the basis of an assay encompassed by the present invention. These and other assays are known in the art and their use with the GFPs and BFPs of the present invention will become obvious to one skilled in the art in light of the teachings disclosed herein. Examples of such assays include competitive assays wherein labeled and unlabeled ligands competitively bind to a ligator, noncompetitive assay where a ligand is captured by a ligator and either measured directly or "sandwiched" with a secondary ligator that is labeled. Still other types of assays include immunoassays, single-step homogeneous assays, multiple-step heterogeneous assays, and enzyme assays.

In a number of embodiments, the mutant GFPs and BFPs are combined with fluorescent microscopy using known techniques (see, e.g., Stauber et al., Virol. 213:439-454 (1995)) or preferably with fluorescence activated cell sorting (FACS) to detect and optionally purify or clone cells that express specific recombinant constructs. For a brief overview of the FACS and its uses, see: Herzenberg et al., 1976, "Fluorescence activated cell sorting", Sci. Amer. 234, 108; see also FLOW CYTOMETRY AND SORTING, eds. Melamad, Mullaney and

Mandelsohn, John Wiley and Sons, Inc., New York, 1979). Briefly, fluorescence activated cell sorters take a suspension of cells and pass them single file into the light path of a laser placed near a detector. The laser usually has a set wavelength. The detector measures the fluorescent emission intensity of each cell as it passes through the instrument and generates a histogram plot of cell number versus fluorescent intensity. Gates or limits can be placed on the histogram thus identifying a particular population of cells. In one embodiment, the cell sorter is set up to select cells having the highest probe intensity, usually a small fraction of the cells in the culture, and to separate these selected cells away from all the other cells. The level of intensity at which the sorter is set and the fraction of cells which is selected, depend on the condition of the parent culture and the criteria of the isolation. In general, the operator should first sort an aliquot of the culture, and record the histogram of intensity versus number of cells. The operator can then set the selection level and isolate an appropriate number of the most active cells. Currently, fluorescence activated cell sorters are equipped with automated cell cloning devices. Such a device enables one to instruct the instrument to singly deposit a selected cell into an individual growth well, where it is allowed to grow into a monoclonal culture. Thus, genetic homogeneity is established within the newly cloned culture.

#### IV. General Applications for the GFP Mutants

It should be self-evident that the mutant GFP and BFP sequences described here have unlimited uses, particularly as signal or reporter sequences for the co-expression of other nucleic acid sequences of interest and/or to track the location and/or movement of other sequences within the cell, within tissue and the like. For example, these reporter type sequences could be used to track the spread (or lack thereof) of a disease causal agent in drug screening assays or could

readily be used in diagnostics. Some of the more interesting applications are described below.

**A. Protein Trafficking**

5        Normally, expressed mutant GFPs and BFPs are distributed throughout the cell (particularly mammalian cells), except for the nucleolus. However, as described below, when a GFP mutant is fused to the HIV-1 Rev protein, a hybrid molecule results which retains the Rev function and is  
10      localized mainly in the nucleolus where Rev is found. Fusion to the N-terminal domain of the HIV-1 Nef protein produces a hybrid protein detectable in the plasma membrane. Thus, the GFP mutants can be used to monitor the subcellular targeting and transport of proteins to which they are fused.

15

**B. Gene Therapy**

The mutant GFPs described here have interesting and useful applications in gene therapy. Gene therapy in general is the correction of genetic defects by insertion of exogenous  
20      cellular genes that encode a desired function into cells that lack that function, such that the expression of the exogenous gene a) corrects a genetic defect or b) causes the destruction of cells that are genetically defective. Methods of gene therapy are well known in the art, see, for example, Lu, M.,  
25      et al. (1994), *Human Gene Therapy* 5:203; Smith, C. (1992), *J. Hematotherapy* 1:155; Cassel, A., et al. (1993), *Exp. Hematol.* 21:585 (1993); Larrick, J.W. and Burck, K.L., *GENE THERAPY: APPLICATION OF MOLECULAR BIOLOGY*, Elsevier Science Publishing Co., Inc., New York, New York (1991) and Kreigler, M. *GENE TRANSFER  
30      AND EXPRESSION: A LABORATORY MANUAL*, W.H. Freeman and Company, New York (1990), each incorporated herein by reference. One modality of gene therapy involves (a) obtaining from a patient a viable sample of primary cells of a particular cell type;  
35      (b) inserting into these primary cells a nucleic acid segment encoding a desired gene product; (c) identifying and isolating cells and cell lines that express the gene product; (d) re-introducing cells that express the gene product; (e) removing from the patient an aliquot of tissue including cells

resulting from step c and their progeny; and (f) determining the quantity of the cells resulting from step c and their progeny, in said aliquot. The introduction into cells in step c of a polycistronic vector that encodes GFP or BFP in addition to the desired gene allows for the quick identification of viable cells that contain and express the desired gene.

Another gene therapy modality involves inserting the desired nucleic acid into selected tissue cells *in situ*, for example into cancerous or diseased cells, by contacting the target cells *in situ* with retroviral vectors that encode the gene product in question. Here, it is important to quickly and reliably assess which and what proportion of cells have been transfected. Co-expression of GFP and BFP permits a quick assessment of proportion of cells that are transfected, and levels of expression.

#### C. Diagnostics

One potential application of the GFP/BFP variants is in diagnostic testing. The GFP/BFP gene, when placed under the control of promoters induced by various agents, can serve as an indicator for these agents. Established cell lines or cells and tissues from transgenic animals carrying GFP/BFP expressed under the desired promoter will become fluorescent in the presence of the inducing agent.

Viral promoters which are transactivated by the corresponding virus, promoters of heat shock genes which are induced by various cellular stresses as well as promoters which are sensitive to organismal responses, e.g. inflammation, can be used in combination with the described GFP/BFP mutants in diagnostics.

In addition, the effect of selected culture conditions and components (salt concentrations, pH, temperature, trans-acting regulatory substances, hormones, cell-cell contacts, ligands of cell surface and internal receptors) can be assessed by incubating cells in which sequences encoding the fluorescent proteins provided herein are operably linked to nucleic acids (especially regulatory

elements such as promoters) derived from a selected gene, and detecting the expression and location of fluorescence.

#### D. Toxicology

Another application of the GFP/BFP-based methodologies is in the area of toxicology. Assessment of the mutagenic potential of any compound is a prerequisite for its use. Until recently, the Ames assay in *Salmonella* and tests based on chromosomal aberrations or sister chromatid exchanges in cultured mammalian cells were the main tools in toxicology. However, both assays are of limited sensitivity and specificity and do not allow studies on mutation induction in various organs or tissues of the intact organism.

The introduction of transgenic mice with a mutational target in a shuttle vector has made possible the detection of induced mutations in different tissues *in vivo*. The assay involves DNA isolation from tissues of exposed mice, packaging of the target DNA into bacteriophage lambda particles and subsequent infection of *E. coli*. The mutational target in this assay is either the lacZ or lacI genes and quantitation of blue vs white plaques on the bacterial lawn allows for mutagenic assessment.

GFP/BFP could significantly simplify both the tissue culture and transgenic mouse procedures. Expression of GFP/BFP under the control of a repressor, which in turn is driven by the promoter of a constitutively expressed gene, will establish a rapid method for evaluating the mutagenic potential of an agent. The presence of fluorescent cells, following exposure of a cell line, tissue or whole animal carrying the GFP/BFP-based detection construct, will reflect the mutagenicity of the compound in question. GFP/BFP expressed under the control of the target DNA, the repressor gene, will only be synthesized when the repressor is inactivated or turned off or the repressor recognition sequences are mutated. Direct visualization of the detector cell line or tissue biopsy can qualitatively assess the mutagenicity of the agent, while FACS of the dissociated cells can provide for quantitative analysis.

#### E. Drug Screening

The GFP/BFP detection system could also significantly expedite and reduce the cost of some current drug screening procedures. A dual color screening system (DCSS), in which GFP is placed under the promoter of a target gene and BFP is expressed from a constitutive promoter, could provide for rapid analysis of agents that specifically affect the target gene. Established cell lines with the DCSS could be screened with hundreds of compounds in few hours. The desired drug will only influence the expression of GFP. Non-specific or cytotoxic effects will be detected by the second marker, BFP. The advantages of this system are that no exogenous substances are required for GFP and BFP detection, the assay can be used with single cells, cell populations, or cell extracts, and that the same detection technology and instrumentation is used for very rapid and non-destructive detection.

The search for antiviral agents which specifically block viral transcription without affecting cellular transcription, could be significantly improved by the DCSS. In the case of HIV, appropriate cell lines expressing GFP under the HIV LTR and BFP under a cellular constitutive promoter, could identify compounds which selectively inhibit HIV transcription. Reduction of only the green but not the blue fluorescent signal will indicate drug specificity for the HIV promoter. Similar approaches could also be designed for other viruses.

Furthermore, the search for antiparasitic agents could also be helped by the DCSS. Established cell lines or transgenic nematodes or even parasitic extracts where expression of GFP depends on parasite-specific trans splicing sequences while BFP is under the control of host-specific cis splicing elements, could provide for rapid screen of selective antiparasitic drugs.

The invention will be more readily understood by reference to the following specific examples which are

included for purposes of illustration only and are not intended to limit the invention unless so stated.

#### **EXAMPLES**

The following general protocol was used to generate mutant GFP- or BFP-encoding nucleic acids, transform host cells, and express the mutant GFP and BFP proteins:

- Clone a nucleic acid that encodes either wtGFP or BFP( $Tyr_{67} \rightarrow His$ ), under the control of eukaryotic or prokaryotic promoters, into a standard ds-DNA plasmid
- Convert the plasmid vector to a ss-DNA by standard methods
- Anneal the ss-DNA to 40-50 nucleotide DNA oligomers having base mismatches at the site(s) intended to be engineered
- Convert the ss-DNA to a closed ds-DNA plasmid vector by use of DNA polymerase and standard protocols
- Identify plasmids containing the desired mutations by restriction analysis following plasmid DNA isolation from *E. coli* strains transformed with the mutagenized DNA
- verify the presence of mutations by DNA sequencing
- transfet human transformed embryonic kidney 293 cells with equal amounts of DNA from the appropriate plasmids
- compare the fluorescence intensity of the signals

#### **Nucleic acids and vectors**

The wtGFP cDNA (SEQ ID NO:1) was obtained from Dr. Chalfie of Columbia University. All mutants described were obtained by modifying this wtGFP sequence as detailed below.

The vectors used to clone and to express the GFPs and BFPs are derivatives of the commercially available plasmids pcDNA3 (Invitrogen, San Diego, CA), pBSSK+ (Stratagene, La Jolla, CA) and pET11a (Novagen, Madison, WI).

wtGFP protein expression in mammalian cells

Several vectors for the expression of GFP in mammalian cells were constructed:

pFRED4 carries the wtGFP sequences under the control of the cytomegalovirus (CMV) early promoter and the polyadenylation signal of the Human Immunodeficiency Virus-1 (HIV) 3' Long Terminal Repeat (LTR). To derive pFRED4 we amplified the GFP coding sequence from plasmid #TU58 (Chalfie et al., 1994) by the polymerase chain reaction (PCR). For PCR amplification of the GFP coding region, oligonucleotides #16417 and #16418 were used as primers. Oligonucleotide #16417:

5'-GGAGGCGCGCAAGAAATGGCTAGCAAAGGAGAAGA-3' (SEQ ID NO:3), containing the BssHII recognition sequence and the translation initiation sequence of the HIV-1 Tat protein, was the sense primer. The antisense primer, #16418:

5'-GCAGGGATCCTTATTGTATAGTTCATCCATGCCATG-3' (SEQ ID NO:4) contained the BamHI recognition sequence. The amplified fragment was digested with BssHII and BamHI and cloned into BssHII and BamHI digested pCMV37M1-10D, a plasmid containing the CMV early promoter and the HIV-1 p37gag region, followed by several cloning sites and the HIV-1 3' LTR. Thus the p37gag gene was replaced by GFP, resulting in pFRED4.

In a second step, the 1485bp fragment from pFRED4, generated from StuI and BamHI double digestion, was subcloned into the 4747bp vector derived from the NruI and BamHI double digestion of pcDNA3. The resulting plasmid, pFRED7 (SEQ ID NO:5), expresses GFP under the control of the early CMV promoter and the bovine growth hormone polyadenylation signal.

Bacterial expression

For bacterial expression, we constructed plasmid pBSGFP (SEQ ID NO:6), a pBSSK+ derivative carrying wtGFP. pBSGFP was generated by inserting the GFP containing region of pFRED4, digested with BamHII and BamHI and subsequently treated with Klenow, into the EcoRV digested pBSSK+ vector. In pBSGFP the wtGFP is fused downstream to the 43 amino acids of the alpha peptide of beta galactosidase, present in the pBSSK+ polylinker region. The added amino acids at the

N-terminus of wtGFP have no apparent effect on the GFP signal, as judged from subsequent plasmids containing precise deletions of the extra amino acids.

For GFP overexpression and purification we generated plasmid pFRED13 (SEQ ID NO:7) by ligating the 717bp fragment from pFRED7 digested with NheI and BamHI, to the 5644bp fragment resulting from the NheI and BamHI double digestion of pET11a. In pFRED13, GFP is synthesized under the control of the bacteriophage T7 ph10 promoter.

The oligonucleotides used for GFP mutagenesis were synthesized by the DNA Support Services of the ABL Basic Research Program of the National Cancer Institute. DNA sequencing was performed by the PCR-assisted fluorescent terminator method (ReadyReaction DyeDeoxy Terminator Cycle Sequencing Kit, ABI, Columbia, MD) according to the manufacturer's instructions. Sequencing reactions were resolved on the ABI Model 373A DNA Sequencing System. Sequencing data were analyzed using the Sequencher program (Gene Codes, Ann Arbor, MI).

Enzymes were purchased from New England Biolabs (Beverly, MA) and used according to conditions described by the supplier. Chemicals used for the purification of wild type and mutant proteins were purchased from SIGMA (St. Louis, MO). Tissue culture media were obtained from Biofluids (Rockville, MD) and GIBCO/BRL (Gaithersburg, MD). Competent bacterial cells were purchased from GIBCO/BRL.

#### Preparation of mutants

Initially, plasmid pBSGFP was used to mutagenize the GFP coding sequence by single-stranded DNA site directed mutagenesis, as described by Schwartz et al. (1992) *J. Virol.* 66:7176. In addition to changing specific codons, our strategy was also to improve GFP expression by replacing potential inhibitory nucleotide sequences without altering the GFP amino acid sequence. This approach has been successfully employed in the past for other proteins (Schwartz et al. (1992) *J. Virol.* 66:7176).

For the pBSGFP mutagenesis the following oligonucleotides were used:

#17422 (SEQ ID NO:8):

5'-CAATTTGTGTCCCAGAATGTTGCCATCTTCCTTGAAGTCAATACCTTT-3'

#17423 (SEQ ID NO:9):

5'-GTCTTGAGTTGCCGTATCTTGAAGAAGATGCTCCTTCCTGTAC-3'

#17424 (SEQ ID NO:10):

5'-CATGGAACAGGCAGTTGCCAGTAGTGCAGATGAACCTCAGGGTAAGTTTC-3'

#17425 (SEQ ID NO:11):

5'-CTCCACTGACAGAGAACCTGTGGCCGTTAACATCACCATC-3'

#17426 (SEQ ID NO:12):

5'-CCATCTTCAATGTTGTGGCGGTCTTGAAGTTCACTTGATTCCATT-3'

#17465 (SEQ ID NO:13):

5'-CGATAAGCTTGAGGGATCCTCAGTTGTACAGTTCATCCATGC-3'

Oligonucleotide #17426 introduces a mutation in GFP, converting the Isoleucine (Ile) at position 168 into Threonine (Thr). The Ile168Thr change has been shown to alter the GFP spectrum and to also increase the intensity of GFP fluorescence by almost two-fold at the emission maxima (Heim et al. (1994), *supra*).

The mutagenesis mixture was used to transform DH5a competent *E. coli* cells. Ampicillin resistant colonies were obtained and examined for their fluorescent properties by excitation with UV light. One colony, significantly brighter than the rest, was apparent on the agar plate. This colony was further purified, the plasmid DNA was isolated and used to transform DH5a competent bacteria. This time all the colonies were bright green when excited with the UV light, indicating that the bright green fluorescence was associated with the presence of the plasmid. The sequence of the GFP segment (SEQ ID NO:14, representing only the segment and not the whole plasmid) of this plasmid, called pBSGFPsg11, was then determined. The sequence analysis revealed that in addition to the designed nucleotide changes, which do no alter the amino acid sequence of GFP, and the Ile168Thr mutation, a second spontaneous mutation had occurred. A thymidine at position 322 of SEQ ID NO:14, which is the GFP-coding region of the pBSGFPsg11 DNA, was replaced by a cytosine. This

nucleotide change converts the phenylalanine (Phe) at position 65 of the GFP amino acid sequence into a leucine (Leu). A series of experiments, which will be described below, demonstrated that indeed the Phe65Leu mutation was responsible for the increase in the intensity of the fluorescent GFP signal.

In subsequent experiments, involving generation of rationally designed GFP mutant combinations to be detailed below, we also used the single-stranded DNA site directed mutagenesis approach. This time, however, the template DNAs were pFRED7 derivatives instead of pBSGFP.

#### Transfection and expression

The 293 cell line, an adenovirus-transformed human embryonal kidney cell line (Graham et al. (1977), *J. Gen. Virol.* 5:59) was used for protein expression analysis. The cells were cultured in Dulbecco's modified culture medium (DMEM) supplemented with 10% heat-inactivated fetal bovine serum (FBS, Biofluids).

Transfection was performed by the calcium phosphate coprecipitation technique as previously described (Graham et al. (1973), *Virol.* 52:456; Felber et al. (1990), *J. Virol.* 64:3734. Plasmid DNA was purified by Qiagen columns according to the manufacturer's instructions (Qiagen). A mix of 5 to 10 µg of total DNA per ml of final precipitate was overlaid on the cells in 60 mm or 6- and 12-well tissue culture plates (Falcon), using 0.5, 0.25 and 0.125 ml of precipitate, respectively. After overnight incubation, the cells were washed, placed in medium without phenol red and measured in a plate spectrofluorometer, e.g., Cytofluor II (Perceptive Biosystems, Framingham, MA.)

#### Purification of wild-type and mutant proteins:

*E. coli* strains carrying pFRED13 or other pET11a derivatives with mutant GFP genes were used for the overproduction and purification of the wt and mutant GFPs or BFPs. The cells were grown in 1 liter LB broth containing 100

$\mu\text{g}/\text{ml}$  ampicillin at 32° C to a density of 0.6-0.8 optical density units at 600 nm. At this point, the cells were induced with 0.6 mM IPTG and incubated for four more hours. Following harvesting of the cell pellets, cellular extracts were prepared as described by Johnson, B.H and Hecht, M.H., 1994, *Biotechnol.* 12: 1357.

GFPs and BFPs were purified from the cellular extracts as follows: Ammonium sulfate (AS) was added first to the extracts (50g AS per 100g supernatant) to precipitate the proteins. The precipitants were collected by centrifugation at 7500 x g for 15 min and the pellets were dissolved in 5ml of 1 M AS. The samples were then loaded on phenylsepharose column (HR10/10, Pharmacia, Piscataway, NJ) and washed with 20 mM 2-[N-morpholino] ethanesulfonic Acid (MES) pH 5.6 and 1 M AS. Proteins were eluted with a 45 ml gradient to 20 mM MES, pH 5.6. Fractions containing the GFP or BFP protein were colored even under visible light.

Green or blue-colored fractions were further purified on Q-sepharose (Mono Q, HR5/5, Pharmacia) with a 20 ml gradient from 20 mM Tris pH 7.0 to 20 mM Tris pH 7.0, 0.25 M NaCl.

The AS precipitation step was performed at 4° C while the chromatographic procedures were performed at room temperature.

#### Determination of protein concentration

Protein concentrations were determined using the commercially available Bradford protein assay (BioRad, Hercules, CA) with bovine IgG protein as a standard.

#### Analytical polyacrylamide gels

Analytical polyacrylamide gel electrophoresis was used to visualize the degree of purity of the purified GFP or BFP proteins. In all cases, 1 mm thick, 12% acrylamide gels (containing 0.1% SDS, in Tris buffer, pH 7.4) were used, and electrophoresis was performed for 2 hours at 120 V. Gels were stained with Coomassie Blue to visualize the proteins.

Fluorescence measurements

Excitation and emission spectra of solutions of the fluorescent proteins were obtained using a Perkin Elmer L550B spectrofluorimeter (Perkin Elmer, Advanced Biosystems, Foster City., CA).

The relative fluorescence data for the GFP mutants in Table I below were obtained by comparing the cellular fluorescence of the GFP mutants expressed in the transformed human embryonic kidney cell line 293 with wtGFP expressed in the same cell line. Likewise, the relative fluorescence data for the BFP mutants in Table I below were obtained by comparing the cellular fluorescence of the BFP mutants expressed in 293 cells with BFP(Tyr<sub>67</sub>-His) expressed in the same cell line. Equal amounts of DNA encoding wild type or mutant proteins were introduced into 293 cells. Cellular fluorescence was quantified 24 h or 48 hr. post-transfection using Cytofluor II.

A list of GFP mutant proteins indicating the introduced amino acid mutations is shown in Table I.

TABLE I: GFP and BFP mutants

PROTEIN	Amino Acid Position					
	65	66	67	164	168	239
wt GFP	F	S	Y	V	I	K
SG12	L					
SG11	L				T	N
SG25	L	C			T	N
BFP			H			
SB42	L		H			
SB49			H	A		
SB50	L		H	A		

Example 1: SG12

A number of the unique mutants described herein derive from the discovery of an unplanned and unexpected mutation called "SG12", obtained in the course of site-

directed mutagenesis experiments, wherein a phenylalanine at position 65 of wtGFP was converted to leucine. SG12 was prepared as follows: Two plasmids carrying SG12 (SEQ ID NO:15) were generated, pFRED12 for expression in mammalian cells, and pFRED16 for expression in *E. coli* and protein purification. pFRED12 was constructed by ligating the 1557 bp fragment from the double digestion of pFRED7 with Avr II and Pml I into the 4681 bp fragment generated from the Avr II and Pml I digestion of pFRED11 (see below). pFRED16 was derived by subcloning the 717bp segment resulting from the digestion of pFRED12 with NheI and BamHI to the 5644bp fragment of the pET11a vector digested with the same restriction enzymes.

The specific activity of SG12 was about 9-12 times that of wtGFP. See Table II.

Example 2: SG11

A mutant referred to as "SG11," which combined the phenylalanine 65 to leucine alteration with an isoleucine 168 to threonine substitution and a lysine 239 to asparagine substitution, gave a further enhanced fluorescence intensity. SG11 was prepared as follows: Two plasmids carrying SG11 (SEQ ID NO:16) were generated: pFRED11 for expression in mammalian cells and pFRED15 for expression in *E. coli* and protein purification. pFRED11 was constructed by ligating the 717bp region from pBSGFPsg11 DNA digested with NheI and BamHI to the 5221bp fragment derived from the digestion of pFRED7 with the same enzymes. pFRED15 was generated by subcloning the 717bp segment resulting from the digestion of pFRED11 with NheI and BamHI to the 5644 bp fragment of the pET11a vector, digested with the same restriction enzymes.

The mutant SG11 encodes an engineered GFP wherein the alteration comprises the conversion of phenylalanine 65 to leucine and the conversion of isoleucine 168 to threonine. The additional alteration of the C-terminal lys 239 to asn is without effect; the C-terminal lys or asn may be deleted without affecting fluorescence. The specific activity of SG11 is about 19-38 times that of wtGFP. See Table II.

Example 3: SG25

A third and further improved GFP mutant was obtained by further mutating "SG11." This mutant is referred to as "SG25" and comprises, in addition to the SG11 substitutions, and additional substitution of a cysteine for the serine normally found at position 66 in the sequence. SG11 was prepared as follows: Two plasmids carrying SG25 (SEQ ID NO:17) were generated: pFRED25 for expression in mammalian cells and pFRED63 for expression in *E. coli* and protein purification. pFRED25 was constructed by site directed mutagenesis of pFRED11, using oligonucleotide #18217 (SEQ ID NO:18): 5'-CATTGAACACCATAGCACAGAGTAGTGAUTGACTAGTGTTGGCC-3'. This oligonucleotide incorporates the Ser66Cys mutation into SG11. Ser66Cys had been shown to both alter the GFP excitation maxima without significant change in the emission spectrum and to also increase the intensity of the fluorescent signal of GFP (Heim et al., 1995).

pFRED63 was generated by subcloning the 717 bp segment resulting from the digestion of pFRED25 with NheI and BamHI to the 5644 bp fragment of the pET11a vector, digested with the same restriction enzymes.

The mutant SG25 encodes an engineered GFP wherein the alteration comprises the conversion of phenylalanine 65 to leu, the conversion of isoleucine 168 to threonine and the conversion of serine 66 to cysteine. As with SG11, the additional alteration of the C-terminal lysine 239 to asparagine is without effect; the C-terminal lysine or asparagine may be deleted without affecting fluorescence. The specific activity of SG25 is about 56 times that of wtGFP. See Table II.

Example 4: Additional green fluorescent mutants

Additional alterations at different amino acids of the wtGFP, when combined with SG11 and SG25, yielded proteins having at least 5X greater cellular fluorescence compared to the wtGFP. A non-limiting list of these mutations is provided below:

**GFP variants with enhanced cellular fluorescence**

<u>Protein</u>	<u>Altered Amino Acids</u>
SG20	F65L, S66T, I168T, K239N
SG21	F65L, S66A, I168T, K239N
SG27	Y40L, F65L, I168T, K239N
SG30	F47L, F65L, I168T, K239N
SG32	F72L, F65L, I168T, K239N
SG43	F65L, I168T, Y201L, K239N
SG46	F65L, V164A, I168T, K239N
SG72	F65L, S66C, V164A, I168T, K239N
SG91	F65L, S66C, F100L, I168T, K239N
SG94	F65L, S66C, Y107L, I168T, K239N
SG95	F65L, S66C, F115L, I168T, K239N
SG96	F65L, S66C, F131L, I168T, K239N
SG98	F65L, S66C, Y146L, I168T, K239N
SG100	F65L, S66C, Y152L, I168T, K239N
SG101	F65L, S66C, I168T, Y183L, K239N
SG102	F65L, S66C, I168T, F224L, K239N
SG103	F65L, S66C, I168T, Y238L, K239N
SG106	F65L, S66T, V164A, I168T, K239N

**Example 5: SB42**

The blue fluorescent proteins described here and below were derived from the known GFP mutant (Heim et al., PNAS, 1994) wherein histidine is substituted for tyrosine at position 67. We have designated this known mutant BFP(Tyr<sub>67</sub>→His). BFP(Tyr<sub>67</sub>→His) has a shifted emission spectrum. It emits blue light, i.e., it is a blue fluorescent protein (BFP).

By introducing the same mutation in BFP(Tyr<sub>67</sub>→His) that was used to generate SG12, i.e., leucine for phenylalanine at position 65, we created a new mutant that has unexpectedly high fluorescence that we refer to as "SuperBlue-42" (SB42). SB42 was prepared as follows: Two plasmids carrying SB42 (SEQ ID NO:19) were generated: pFRED42 for expression in mammalian cells and pFRED65 for expression in *E. coli* and protein purification. pFRED42 was constructed by site directed mutagenesis of pFRED12, using oligonucleotide

#bio25 (5'-CATTGAAACACCATGAGAGAGACTAGTGACTAGTGTGGCC-3') (SEQ ID NO:20). This oligonucleotide incorporates the Tyr<sub>67</sub>→His mutation into SG12, thus generating the Phe65Leu, Tyr<sub>67</sub>→His double mutant.

pFRED65 was created by subcloning the 717 bp segment resulting from the digestion of pFRED42 with NheI and BamHI to the 5644 bp fragment of the pET11a vector, digested with the same restriction enzymes.

The mutant SB42 encodes an engineered BFP wherein the alterations comprise the conversion of tyrosine 67 to histidine and the conversion of phenylalanine 65 to leucine. The specific activity of SB42 is about 27 times that of BFP(Tyr<sub>67</sub>→His). See Table II.

**Example 6: SB49**

An independent mutation of BFP(Tyr<sub>67</sub>→His) which substitutes the valine at position 164 with an alanine is referred to as "SB49." SB49 was prepared as follows: Plasmid pFRED49 expresses SB49 (SEQ ID NO:21) in mammalian cells. pFRED49 was generated by site directed mutagenesis of pFRED12, using oligonucleotides #19059 and #bio24. Oligonucleotide #19059 (5'-CTTCAATGTTGGCGGATCTTGAAGTTCGCTTGATTCCATTC-3') (SEQ ID NO:22) introduces the Val164Ala mutation in SG12 while oligonucleotide #bio24 (5'-CATTGAAACACCATGAGAGAAAGTAGTGACTAGTGTGGCC-3') (SEQ ID NO:23) reverts the Phe65Leu alteration to the wt sequence and, at the same time, incorporates the Tyr<sub>67</sub>→His mutation.

The mutant SB49 encodes an engineered BFP wherein the alterations comprise the conversion of tyrosine 67 to histidine, and the conversion of valine 164 to alanine. The specific activity of SB49 was about 37 times that of BFP(Tyr<sub>67</sub>→His). See Table II.

**Example 7: SB50**

A combination of the above two BFP mutations resulted in "SB50," which gave an even greater fluorescence enhancement than either of the previous mutations. SB50 was prepared as follows: Two plasmids carrying SB50 (SEQ ID NO:

24) were generated: pFRED50 for expression in mammalian cells and pFRED67 for expression in *E. coli* and protein purification. pFRED50 was constructed by site directed mutagenesis of pFRED12, using oligonucleotides #19059 and #bio25.

pFRED67 was created by subcloning the 717bp segment resulting from the digestion of pFRED50 with NheI and BamHI to the 5644 bp fragment of the pET11a vector digested with the same restriction enzymes.

The mutant SB50 encodes an engineered BFP wherein the alterations comprise the conversion of tyrosine 67 to histidine, the conversion of phenylalanine 65 to leucine and the conversion of alanine 164 to valine. The specific activity of SB50 was about 63 times that of BFP(Tyr<sub>67</sub>→His). See Table II.

TABLE II

Mutant	Excitation Maximum (nm)	Emission Maximum (nm)	Factor of increased green fluorescence (at maximum emission) as compared to wtGFP	Factor of increased blue fluorescence (at maximum emission) as compared to BFP(Tyr <sub>67</sub> →His)
SG12	398	509	9-12X	
SG11	471	508	19-38X	
SG25	473	509	50-100X	
SB42	387	450		27X
SB49	387	450		37X
SB50	387	450		63X

The dramatic increase in fluorescent activity resulting from the amino acid substitutions of the present invention was wholly unexpected. The cellular fluorescence of the mutants was at least five times greater, and usually over twenty times greater, than that of the parent wtGFP or BFP(Tyr<sub>67</sub>→His). Note that the maximum emission wavelengths vary among the mutants, and that the above-reported fold

increases refer only to minimal increases in relative cellular fluorescence at the maximum emission wavelength of the mutant. Given a particular wavelength, the values may be substantially larger, i.e., the mutants may have a 200-fold greater cellular fluorescence than the reference wtGTP or BFP(Tyr<sub>67</sub>>His). This is important because devices for measuring fluorescence often have set wavelengths, or the limitations of a given experiment often require the use of a set wavelength. Thus, for example, the emission and detection parameters of a fluorescence microscope or a fluorescence-activated cell sorter may be set for a wavelength wherein the cellular fluorescence of a given mutant is 200-fold greater than that of the known GFPs and BFPs.

The GFP and BFP mutants of this invention, in contrast to the wild type protein or other reported mutants, allow detection of green fluorescence in living mammalian cells when present in few copies stably integrated into the genome. This high cellular fluorescence of the mutant GFPs and BFPs is useful for rapid and simple detection of gene expression in living cells and tissues and for repeated analysis of gene expression over time under a variety of conditions. They are also useful for the construction of stable marked cell lines that can be quickly identified by fluorescence microscopy or fluorescence activated cell sorting.

Example 8

We have established fluoroplate-based assays for the quantitation of gene expression after transfections. In a number of embodiments, a nucleic acid encoding a mutant GFP or BFP of this invention is inserted into a vector and introduced into and expressed in a cell. Typically, expression of GFP mutants can be detected as quickly as 5 hours post-infection or less. Expression is followed over time in living cells by a simple measurement in multi-well plates. In this way, many transfections can be processed in parallel.

Example 9

The vectors and nucleic acids provided herein are used to generate chimeric proteins wherein a nucleic acid sequence that encodes a selected gene product is fused to the 5 C- or N-terminus of the mutant GFPs and/or BFPs of this invention. A number of unique viral, plasmid and hybrid gene constructs have been generated that incorporate the new mutant GFP and/or mutant BFP sequences indicated above. These include:

10 • HIV viral sequences (in the nef gene) containing SG11 or SG25

• Neomycin & hygromycin plasmids containing SG11 or SG25

• Moloney Leukemia Virus vector (retrovirus) also expressing SG25

15 • Hybrid gene constructs expressing HIV viral proteins (rev, td-rev, tat, nef, gag, env, and vpr) and either SG11 or SG25 or SB50.

• Hybrid gene construct containing vectors that incorporate the cytoplasmic proteins ran, B23, nucleolin, poly-A 20 binding protein and either SG11 or SG25 or SB50.

These hybrids of the mutant nucleic acids provided herein are used to study protein trafficking in living mammalian cells. Like the wild type GFP, the mutant GFP 25 proteins are normally distributed throughout the cell except for the nucleolus. Fusions to other proteins redistribute the fluorescence, depending on the partner in the hybrid. For example, fusion with the entire HIV-1 Rev protein results in a hybrid molecule which retains the Rev function and is 30 localized in the nucleolus where Rev is preferentially found. Fusion to the N-terminal domain of the HIV-1 Nef protein created a chimeric protein detected in the plasma membrane, the site of Nef localization.

35 Example 10: pCMVgfol1

pCMVgfol1 is a pFRED11 derivative containing the bacterial neomycin phosphotransferase gene (neo) (Southern and Berg (1982) *J. Mol. Appl. Genetics* 1:327) fused at the

C-terminus of SG11. A four amino acid (Gly-Ala-Gly-Ala) (SEQ ID NO:26) linker region connects the last amino acid of SG11 to the second amino acid of neo, thus generating the hybrid SG11-neo protein (gfo11, SEQ ID NO:25). Gfo11 is expressed from the CMV promoter and contains the intact SG11 polypeptide and all of neo except for the first Met.

5 pCMVgfo11 was constructed in several steps. First, pFRED11DNae was constructed by NaeI digestion of pFRED11 and self-ligation of the 4613bp fragment. The NaeI deletion removes the SV40 promoter and neo gene from pFRED11, thus creating pFRED11DNae. Next, in order to fuse the neo coding region downstream to SG11, the neo gene was PCR amplified from pcDNA3 using primers Bio51 (5'-CGCGGATCCTTCGAACAAAGATGGATTGCACGC-3') (SEQ ID NO:27) and 10 Bio52 (5'-CCGGAATTCTCAGAAGAACTCGTCAAGAAGGCGA-3') (SEQ ID NO:28). Primer Bio51 introduces a BamHI site followed by a BstBI recognition sequence at the 5' end of neo, while primer Bio52 introduces an EcoRI site 3' to the neo gene. The PCR product was digested with BamHI and EcoRI and cloned into the 15 4582 bp vector resulting from the BamHI-EcoRI digestion of pFRED11DNae, thus generating pFRED11DNaeBstNeo. Subsequently, SG11 was PCR amplified from pFRED11DNae using primers Bio49 (5'-GGCGCGCAAGAAATGGCTAGCAAAGGAGAAGAACTCTTCACTGGAG-3') (SEQ ID NO:29) and Bio50 (5'-CCCATCGATAGCACCGACCAGCACCAGTTGTACAGTTCATCCATGCCATGT-3') (SEQ ID 20 NO:30) to remove the sgII stop codon in pFRED11DNaeBstNeo and to introduce the four amino acid (Gly-Ala-Gly-Ala) linker followed by a ClaI site. The PCR product was digested with NheI and ClaI and cloned into the 4763 bp NhelBstBi fragment 25 from pFRED11DNaeBstNeo, thus generating pCMVgfo11.

Following transfection of 293 cells (Graham et al. (1977), J. Gen. Virol. 5:59) as well as other human and mouse cell lines with pCMVgfo11, bright fluorescent transfected 30 colonies were apparent under the fluorescent microscope and colonies resistant to G418 could be obtained two weeks later.

35 It should be noted that pCMVgfo11 was the best protein fusion in terms of fluorescent emission intensity and

number of G418 resistant colonies compared to several SG11-neo or neo-SG11 fusions generated and examined.

**Example 11: pPGKgfo25**

5 pPGKgfo25 is a pCMVgfoII derivative containing SG25 instead of SG11 within gfo (SEQ ID NO: 31). Expression of gfo25 in pPGKgfo25 is under the control of the mouse phosphoglycerate kinase-1 (PGK) promoter.

10 pPGKgfo25 was constructed in several steps. First, a SacII site was introduced downstream of the PGK promoter in pPGKneobpA (Soriano et al. (1991) Cell: 64-393) by:

15 i) annealing oligonucleotides #18990 (SEQ ID NO:32) (5'-GACCGGGACACGTATCCAGCCTCCGC-3') and 18991 (SEQ ID NO:33) (5'-GGAGGCTGGATACGTGTCCCCGGTCTGCA-3') to create a double stranded adapter for PstI at the 5' end and SacII at the 3' end.

20 ii) ligating this adapter to the 3423bp fragment from the PstI-SacII double digestion of pPGKneobpA, thus generating pPGKPtAfSc.

25 Next, the CMV promoter of pFRED25 was replaced with the PGK promoter by cloning the 565bp SaII (filled with Klenow)-SacII region from pPGKPtAfSc to the 5288bp BgIII (filled with Klenow)-SacII fragment from pFRED25, resulting in pFRED25PGK. In the final step, pPGKgfo25 was constructed by ligating the 813bp BgIII-NdeI fragment from pFRED25PGK containing the PGK promoter and SG25, to the 4185bp BgIII-NdeI fragment of pCMVgfo11.

**Example 12: pGen-PGKgfo25RO (SEQ ID NO: 34)**

30 pGen-PGKgfo25RO is a pGen- (Soriano et al. (1991), J. Virol. 65:2314) derivative containing the gfo25 hybrid under the control of PGK promoter. It was constructed by subcloning the 2810bp SaII fragment of pPGKgfo25 into the XhoI site of pGen. In viruses generated from pGen-PGKgfo25RO (see below) 35 transcription originated from the PGK promoter is in reverse orientation (RO) to that initiated from the viral long terminal repeats (LTR).

To generate ecotropic or pseudotyped viruses, pGen-PGKgfo25R0 was co-transfected into 293 cells together with pHIT60 and pHIT123 DNAs (production of ecotropic virus) or with pHIT60 and pHCMV-G DNAs (production of pseudotyped virus). pHIT60 and pHIT123 contain the gag-pol and env coding regions from the Moloney murine leukemia virus (Mo-MLV) respectively, under the control of the CMV promoter (Soneoka et al. (1995), *Nuc. Acid Res.* 23:628. pHCMV-G contains the coding region of the G protein from the vesicular stomatitis virus (VSV) expressed from the CMV promoter (Yee et al. (1994), *Proc. Nat'l Acad. Sci. USA* 91:9564. Virus-containing supernatants were harvested 48 hours post transfection, filtered and stored at -80°C.

15     Example 13: pNLnSG11 (SEQ ID NO:35)

The SG11 sequence from plasmid pFRED11 was PCR-amplified with primers #17982 (SEQ ID NO:36) (5'-GGGGCGTACGGAGCGCTCCGAATCGGTACCGTTAACGGGCCCTCTCGAGTCC GTTGTACAGTTCATCCATG-3') and #17983 (SEQ ID NO:37) (5'-GGGGGAATTCGCGCGTACGTAAGCGCTAGCTGAGCAAGAAATGGCTAGCAAA GGAGAAGAACTC-3'). The PCR product was digested with BpI and XhoI and cloned into the large BpI-XhoI fragment from pNL4-3 (Adachi et al. (1986), *J. Virol.* 59: 284. In pNLnSG11 the full SG11 polypeptide containing an additional four linker-encoded amino acids at the C-terminus, is expressed as a hybrid protein with the 24 N-terminal amino acids of the HIV-1 protein Nef.

We constructed transmissible HIV-1 stocks with our mutants, which generate green fluorescence upon transfection of human cells. These transmissible HIV-1 stocks are used to detect the kinetics of infection under a variety of conditions. In particular, they are used to study the effects of drugs on the kinetics of infection. The level of fluorescence, and the subcellular compartmentalization of that fluorescence, is easily visualized and quantified using well known methods. This system is easy to visualize, and dramatically cuts the costs of many experiments that are presently tedious and expensive.

To produce infectious virus, pNLnSG11 was transfected in 293 cells. 24 hours later, Jurkat cells were added to the transfectants. At various times post-infection, the medium was removed, filtered, and used to infect fresh 5 Jurkat or other HIV-1-permissive cells. Two days later the infected cells were green under fluorescent microscope. Visible syncytia were also green. Viral stocks were generated and kept at -80° C.

10 When the nucleic acids, vectors, mutant proteins provided herein are combined with the knowledge of those skilled in the art of genetic engineering and the guidance provided herein, it will be apparent to one of ordinary skill in the art that many changes and modifications can be made 15 thereto without departing from the spirit or scope of the invention as set forth herein. These changes and modifications are encompassed by the present invention.

## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

5

(i) APPLICANT: Pavlakis, George N.  
Gaitanaris, George A.  
Stauber, Roland H.  
Vournakis, John N.

10

(ii) TITLE OF INVENTION: Mutant Aequorea victoria Fluorescent  
Proteins Having Increased Cellular Fluorescence

15

(iii) NUMBER OF SEQUENCES: 37

15

(iv) CORRESPONDENCE ADDRESS:

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20 (D) STATE: California  
(E) COUNTRY: USA  
(F) ZIP: 94111-3834

25

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk  
(B) COMPUTER: IBM PC compatible  
(C) OPERATING SYSTEM: PC-DOS/MS-DOS  
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30

30

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: US Not yet assigned  
(B) FILING DATE: Not yet assigned  
(C) CLASSIFICATION:

35

(viii) ATTORNEY/AGENT INFORMATION:

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40

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45

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 720 base pairs  
(B) TYPE: nucleic acid  
50 (C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

55

(ix) FEATURE:

(A) NAME/KEY: CDS  
(B) LOCATION: 1..720  
60 (D) OTHER INFORMATION: /product= "wild type Aequorea victoria  
Green Fluorescent Protein (wtGF)"

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	GTT GAA TTA GAT GGT GAT GTT AAT GGG CAC AAA TTT TCT GTC AGT GGA	96
	Val Glu Leu Asp Gly Asp Val Asn Gly His Lys Phe Ser Val Ser Gly	
10	20 25 30	
	GAG GGT GAA GGT GAT GCA ACA TAC GGA AAA CTT ACC CTT AAA TTT ATT	144
	Glu Gly Glu Gly Asp Ala Thr Tyr Gly Lys Leu Thr Leu Lys Phe Ile	
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	Cys Thr Thr Gly Lys Leu Pro Val Pro Trp Pro Thr Leu Val Thr Thr	
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	65 70 75 80	
25	CGG CAT GAC TTT TTC AAG AGT GCC ATG CCC GAA GGT TAT GTA CAG GAA	288
	Arg His Asp Phe Phe Lys Ser Ala Met Pro Glu Gly Tyr Val Gln Glu	
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	Arg Thr Ile Phe Phe Lys Asp Asp Gly Asn Tyr Lys Thr Arg Ala Glu	
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	Val Lys Phe Glu Gly Asp Thr Leu Val Asn Arg Ile Glu Leu Lys Gly	
	115 120 125	
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	Ile Asp Phe Lys Glu Asp Gly Asn Ile Leu Gly His Lys Leu Glu Tyr	
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40	AAC TAT AAC TCA CAC AAT GTA TAC ATC ATG GCA GAC AAA CAA AAG AAT	480
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	Gly Ile Lys Val Asn Phe Lys Ile Arg His Asn Ile Glu Asp Gly Ser	
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	Val Gln Leu Ala Asp His Tyr Gln Gln Asn Thr Pro Ile Gly Asp Gly	
	180 185 190	
	CCT GTC CTT TTA CCA GAC AAC CAT TAC CTG TCC ACA CAA TCT GCC CTT	624
	Pro Val Leu Leu Pro Asp Asn His Tyr Leu Ser Thr Gln Ser Ala Leu	
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                   (A) LENGTH: 239 amino acids  
                   (B) TYPE: amino acid  
                   (D) TOPOLOGY: linear

10                  (ii) MOLECULE TYPE: protein

10                  (xii) SEQUENCE DESCRIPTION: SEQ ID NO:2:

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60	Gly	Ile	Lys	Val	Asn	Phe	Lys	Ile	Arg	His	Asn	Ile	Glu	Asp	Gly	Ser
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65	Val	Gln	Leu	Ala	Asp	His	Tyr	Gln	Gln	Asn	Thr	Pro	Ile	Gly	Asp	Gly
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70	Pro	Val	Leu	Leu	Pro	Asp	Asn	His	Tyr	Leu	Ser	Thr	Gln	Ser	Ala	Leu
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                   (A) LENGTH: 35 base pairs  
                   (B) TYPE: nucleic acid  
                   (C) STRANDEDNESS: single  
                   (D) TOPOLOGY: linear

65                  (ii) MOLECULE TYPE: DNA

(ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..35  
 (D) OTHER INFORMATION: /note= "oligonucleotide sense primer  
 #16417"

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

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10

35

(2) INFORMATION FOR SEQ ID NO:4:

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(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 36 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: DNA

25

(ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..36  
 (D) OTHER INFORMATION: /note= "oligonucleotide antisense primer  
 #16418"

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

GCGGGATCCT TATTTGTATA GTTCATCCAT GCCATG

36

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(2) INFORMATION FOR SEQ ID NO:5:

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(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 6238 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

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(ii) MOLECULE TYPE: DNA

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(ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..6238  
 (D) OTHER INFORMATION: /note= "pFRED7"

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

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GACGGATCGG GAGATCTCCC GATCCCCAT GGTCGACTCT CAGTACAATC TGCTCTGATG

60

CCGCATAGTT AAGCCAGTAT CTGCTCCCTG CTTGTGTGTT GGAGGTCGCT GAGTAGTGCG

120

CGAGCAAAT TTAAGCTACA ACAAGGCAAG GCTTGACCGA CAATTGCATG AAGAATCTGC

180

60

TTAGGGTTAG CGGTTTGCGC CTGCTTCGCC TCGAGGCCTG GCCATTGCAT ACGTTGTATC

240

CATATCATAA TATGTACATT TATATTGGCT CATGTCCAAC ATTACCGCCA TGTTGACATT

300

65

GATTATTGAC TAGTTATTAA TAGTAATCAA TTACGGGTC ATTAGTTCAT AGCCCATATA

360

TGGAGTTCCG CGTTACATAA CTTACGGTAA ATGGCCCGCC TGGCTGACCG CCCAACGACC

420

CCCCCCCCATT GACGTCAATA ATGACGTATG TTCCCATAGT AACGCCAATA GGGACTTTCC

480

	ATTGACGTCA ATGGGTGGAG TATTTACGGT AACTGCCCA CTTGGCAGTA CATCAAGTGT	540
	ATCATATGCC AAGTACGCC CCTATTGACG TCAATGACGG TAAATGGCCC GCCTGGCATT	600
5	ATGCCAGTA CATGACCTTA TGGGACTTTC CTACTTGGCA GTACATCTAC GTATTAGTCA	660
	TCGCTATTAC CATGGTGATG CGGTTTGGC AGTACATCAA TGGGCGTGGA TAGCGGTTG	720
	ACTCACGGGG ATTCCAAGT CTCCACCCCA TTGACGTAA TGGGAGTTG TTTTGGCACC	780
10	AAAATCAACG GGACTTTCCA AAATGTCGTA ACAACTCCGC CCCATTGACG CAAATGGCG	840
	GTAGGCGTGT ACGGTGGGAG GTCTATATAA GCAGAGCTCG TTTAGTGAAAC CGTCAGATCG	900
15	CCTGGAGACG CCATCCACGC TGTTTGACC TCCATAGAAC ACACCGGGAC CGATCCAGCC	960
	TCCGCGGGCG CGCAAGAAAT GGCTAGCAAA GGAGAAGAAC TCTTCACTGG AGTTGTCCC	1020
20	ATTCTTGTG AATTAGATGG TGATGTTAAT GGGCACAAAT TTTCTGTCAAG TGGAGAGGGT	1080
	GAAGGTGATG CAACATACGG AAAACTTACC CTTAAATTAA TTTGCACTAC TGGAAAAC	1140
	CCTGTTCCAT GGCCAACACT TGTCACTACT TTCTCTTATG GTGTTCAATG CTTTCAAGA	1200
25	TACCCGGATC ATATGAAACG GCATGACTTT TTCAAGAGTG CCATGCCCGA AGGTTATGTA	1260
	CAGGAAAGAA CTATATTTT CAAAGATGAC GGGAACTACA AGACACGTGC TGAAGTCAAG	1320
	TTTGAAGGTG ATACCCCTGT TAATAGAAC GAGTTAAAAG GTATTGATT TAAAGAAGAT	1380
30	GGAAACATTC TTGGACACAA ATTGGAATAC AACTATAACT CACACAATGT ATACATCATG	1440
	GCAGACAAAC AAAAGAATGG AATCAAAGTT AACTTCAAAA TTAGACACAA CATTGAAGAT	1500
35	GGAAGCGTTC AACTAGCAGA CCATTATCAA CAAAATACTC CAATTGGCGA TGGCCCTGTC	1560
	CTTTTACCAAG ACAACCATTAA CCTGTCCACA CAATCTGCC TTTGAAAGA TCCCAACGAA	1620
40	AAGAGAGACC ACATGGTCT TCTTGAGTTT GTAACAGCTG CTGGGATTAC ACATGGCATG	1680
	GATGAACTAT ACAAAATAAGG ATCCACTAGT AACGGCCGCC AGTGTGCTGG AATTCTGCAG	1740
	ATATCCATCA CACTGGCGGC CGCTCGAGCA TGCACTAGA GGGCCCTATT CTATAGTGT	1800
45	ACCTAAATGC TAGAGCTCGC TGATCAGCCT CGACTGTGCC TTCTAGTTGC CAGCCATCTG	1860
	TTGTTTGCCC CTCCCCCGTG CCTTCCTTGA CCCTGGAAGG TGCCACTCCC ACTGTCCTTT	1920
	CCTAATAAAA TGAGGAAATT GCATCGCATT GTCTGAGTAG GTGTCATTCT ATTCTGGGG	1980
50	GTGGGGTGGG GCAGGACAGC AAGGGGGAGG ATTGGGAAGA CAATAGCAGG CATGCTGGGG	2040
	ATGCGGTGGG CTCTATGGCT TCTGAGGCCG AAAGAACAG CTGGGGCTCT AGGGGGTATC	2100
55	CCCACCGGCC CTGTAGCGGC GCATTAAGCG CGGCGGGTGT GGTGGTTACG CGCAGCGTGA	2160
	CCGCTACACT TGCCAGCGCC CTAGCGCCCG CTCTTCTCGC TTTCTTCCCT TCCTTCTCG	2220
60	CCACGTTCGC CGGCTTCCC CGTCAAGCTC TAAATCGGGG CATCCCTTA GGGTCCGAT	2280
	TTAGTGCTTT ACGGCACCTC GACCCAAAAA AACTTGATTA GGGTGATGGT TCACGTAGTG	2340
	GGCCATCGCC CTGATAGAGCG GTTTTCGCC CTTTGACGTT GGAGTCCACG TTCTTTAATA	2400
65	GTGGACTCTT GTTCCAAACT GGAACAAACAC TCAACCCAT CTCGGTCTAT TCTTTTGATT	2460
	TATAAGGGAT TTTGGGGATT TCGGCCTATT GGTAAAAAAA TGAGCTGATT TAACAAAAAT	2520
	TTAACCGCAA TTAATTCTGT GGAATGTGTG TCAGTTAGGG TGTGGAAAGT CCCCAGGCTC	2580

	CCCAGGCAGG CAGAAGTATG CAAAGCATGC ATCTCAATT A	GTCAGCAACC AGGTGTGGAA	2640
	AGTCCCCAGG CTCCCCAGCA GGCAGAAGTA TGCAAAGCAT GCATCTCAAT TAGTCAGCAA		2700
5	CCATAGTCCC GCCCTTAAC CCGCCCATCC CGCCCTAAC TCCGCCAGT TCCGCCATT		2760
	CTCCGCCCCA TGGCTGACTA ATTTTTTTA TTTATGCAGA GGCGAGGCC GCCTCTGCCT		2820
10	CTGAGCTATT CCAGAAGTAG TGAGGAGGCT TTTTGAGG CCTAGGCTTT TGCAAAAAGC		2880
	TCCCAGGAGC TTGTATATCC ATTTTCGGAT CTGATCAAGA GACAGGATGA GGATCGTTTC		2940
	GCATGATTGA ACAAGATGGA TTGCACGCAG GTTCTCCGGC CGCTTGGGTG GAGAGGCTAT		3000
15	TCGGCTATGA CTGGGCACAA CAGACAATCG GCTGCTCTGA TGCCGCCGTG TTCCGGCTGT		3060
	CAGCGCAGGG GCGCCCGGTT CTTTTGTCA AGACCGACCT GTCCGGTGCC CTGAATGAAC		3120
20	TGCAGGACGA GGCAGCGCGG CTATCGTGC TGGCCACGAC GGGCGTTCT TGCGCAGCTG		3180
	TGCTCGACGT TGTCACTGAA GCGGGAAAGG ACTGGCTGCT ATTGGGCGAA GTGCCGGGGC		3240
	AGGATCTCCT GTCATCTCAC CTTGCTCCTG CCGAGAAAGT ATCCATCATG GCTGATGCAA		3300
25	TGCGCGGCT GCATACGCTT GATCCGGCTA CCTGCCATT CGACCACCAA GCGAACATC		3360
	GCATCGAGCG AGCACGTACT CGGATGGAAG CCGGTCTTGT CGATCAGGAT GATCTGGACG		3420
30	AAGAGCATCA GGGGCTCGCG CCAGCCGAAC TGTTCGCCAG GCTCAAGGCC CGCATGCCCG		3480
	ACGGCGAGGA TCTCGTCGTG ACCCATGGCG ATGCCCTGCTT GCCGAATATC ATGGTGGAAA		3540
	ATGGCCGCTT TTCTGGATTTC ATCGACTGTG GCCGGCTGGG TGTGGCGGAC CGCTATCAGG		3600
35	ACATAGCGTT GGCTACCCGT GATATTGCTG AAGAGCTTGG CGGCGAATGG GCTGACCGCT		3660
	TCCTCGTGCT TTACGGTATC GCCGCTCCCG ATTGCGAGCG CATGCCTTC TATGCCCTTC		3720
40	TTGACGAGTT CTTCTGAGCG GGACTCTGGG GTTCGAAATG ACCGACCAAG CGACGCCAA		3780
	CCTGCCATCA CGAGATTTCG ATTCCACCGC CGCCTTCTAT GAAAGGTTGG GCTTCGGAAT		3840
	CGTTTCCGG GACGCCGGCT GGATGATCCT CCAGCGCGGG GATCTCATGC TGGAGTTCTT		3900
45	CGCCCACCCCC AACITGTTTA TTGCAGCTTA TAATGGTTAC AAATAAAGCA ATAGCATCAC		3960
	AAATTCACA AAATAAGCAT TTTTTCACT GCATTCTAGT TGTGGTTTGT CCAAACATCAT		4020
	CAATGTATCT TATCATGTCT GTATACCGTC GACCTCTAGC TAGAGCTTGG CGTAATCATG		4080
50	GTCATAGCTG TTTCTGTGT GAAATTGTTA TCCGCTACA ATTCCACACA ACATACGAGC		4140
	CGGAAGCATA AAGTGTAAAG CCTGGGGTGC CTAATGAGTG AGCTAACTCA CATTATTGC		4200
55	GTTGCGCTCA CTGCCCGCTT TCCAGTCGGG AAACCTGTGC TGCCAGCTGC ATTAATGAAT		4260
	CGGCCAACGC CGGGGGAGAG CGGGTTGCG TATTGGCGC TCTTCCGCTT CCTCGCTCAC		4320
60	TGACTCGCTG CGCTCGGTGC TTCGGCTGCG GCGAGCGGT A	TCAGCTCACT CAAAGCGGT	4380
	AATAACGGTTA TCCACAGAAAT CAGGGGATAA CGCAGGAAAG AACATGTGAG CAAAAGGCCA		4440
	GCAAAAGGCC AGGAACCGTA AAAAGGCCGC GTTGTGGCG TTTTTCCATA GGCTCCGCC		4500
65	CCCTGACGAG CATCACAAAA ATCGACGCTC AAGTCAGAGG TGGCGAAACC CGACAGGACT		4560
	ATAAAGATAAC CAGGCCTTTC CCCCTGGAAG CTCCCTCGTG CGCTCTCCTG TTCCGACCCCT		4620
	GCCGCTTACC GGATACCTGT CCGCCTTTCT CCCTTCGGGA AGCGTGGCGC TTTCTCAATG		4680

	CTCACGCTGT AGGTATCTCA GTTCGGTGTAA GGTCGTTCGC TCCAAGCTGG GCTGTGTGCA	4740
	CGAACCCCCC GTTCAGCCCG ACCGCTGCCTA CTTATCCGGT AACTATCGTC TTGAGTCCAA	4800
5	CCCGGTAAGA CACGACTTAT CGCCACTGGC AGCAGCCACT GGTAACAGGA TTAGCAGAGC	4860
	GAGGTATGTA GGCAGGTGCTA CAGAGTTCTT GAAGTGGTGG CCTAACTACG GCTACACTAG	4920
10	AAGGACAGTA TTTGGTATCT GCGCTCTGCT GAAGCCAGTT ACCTTCGGAA AAAGAGTTGG	4980
	TAGCTCTGTA TCCGGCAAAAC AAACCACCGC TGGTAGCGGT GGTTTTTTTG TTTGCAAGCA	5040
	GCAGATTACG CGCAGAAAAA AAGGATCTCA AGAAGATCCT TTGATCTTT CTACGGGGTC	5100
15	TGACCGCTCAG TGGAACGAAA ACTCACGTTA AGGGATTTG GTCATGAGAT TATCAAAAAG	5160
	GATCTTCACC TAGATCCTTT TAAATTTAAA ATGAAGTTTT AAATCAATCT AAAGTATATA	5220
20	TGAGTAAACT TGGTCTGACA GTTACCAATG CTTAACAGT GAGGCACCTA TCTCAGCGAT	5280
	CTGTCTATTT CGTTCATCCA TAGTTGCCG ACTCCCCGTC GTGTAGATAA CTACGATACG	5340
	GGAGGGCTTA CCATCTGGCC CCAGTGTGCA AATGATACCG CGAGACCCAC GCTCACCGGC	5400
25	TCCAGATTTA TCAGCAATAA ACCAGCCAGC CGGAAGGGCC GAGCGCAGAA GTGGTCCTGC	5460
	AACTTTATCC GCCTCCATCC AGTCTATTAA TTGTTGCCGG GAAGCTAGAG TAAGTAGTTC	5520
30	GCCAGTTAAT AGTTTGCAGCA ACGTTGTTGC CATTGCTACA GGCATCGTGG TGTCACTGCTC	5580
	GTCGTTGGT ATGGCTTCAT TCAGCTCCGG TTCCCAACGA TCAAGGCGAG TTACATGATC	5640
	CCCCATGTTG TGCAAAAAAAAG CGGTTAGCTC CTTCCGGCCT CCGATCGTTG TCAGAAGTAA	5700
35	GTTGGCCGCA GTGTTATCAC TCATGGTTAT GGCAGCACTG CATAATTCTC TTACTGTCAT	5760
	GCCATCCGTA AGATGCTTTT CTGTGACTGG TGAGTACTCA ACCAAGTCAT TCTGAGAATA	5820
40	GTGTATGCGG CGACCGAGTT GCTCTGCCG GCGCTCAATA CGGGATAATA CGCGGCCACA	5880
	TAGCAGAACT TTAAAAGTGC TCATCATTGG AAAACGTTCT TCGGGGCGAA AACTCTCAAG	5940
	GATCTTACCG CTGTTGAGAT CCAGTTCGAT CTAACCCACT CGTGCACCCA ACTGATCTTC	6000
45	AGCATTCTTT ACTTTCACCA GCGTTCTGG GTGAGCAAA ACAGGAAGGC AAAATGCCGC	6060
	AAAAAAGGGA ATAAGGGCGA CACGGAAATG TTGAATACTC ATACTCTTCC TTTTCAATA	6120
50	TTATTGAAGC ATTATCAGG GTTATTGTCT CATGAGCGGA TACATATTG AATGTATTAA	6180
	GAAAATAAA CAAATAGGGG TTCCGCGCAC ATTTCCCCGA AAAGTGCCAC CTGACGTC	6238

## (2) INFORMATION FOR SEQ ID NO:6:

55           (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3699 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

60           (ii) MOLECULE TYPE: DNA

65           (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..3699
- (D) OTHER INFORMATION: /note= "pBSGFP"

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

	GGAAATTGTA AACGTTAATA TTTTGTAAA ATTGCGTTA AATTTTGTT AAATCAGCTC	60
5	ATTTTTAAC CAATAGGCCG AAATCGCAA AATCCCTTAT AAATCAAAAG AATAGACCGA	120
	GATAGGGTTG AGTGTGTTTC CAGTTGGAA CAAGAGTCCA CTATTAAGA ACGTGGACTC	180
10	CAACGTCAAA GGGCGAAAAA CCGTCTATCA GGGCGATGGC CCACTACGTG AACCATCAC	240
	CTAATCAAGT TTTTGGGGT CGAGGTGCCG TAAAGCACTA AATCGGAACC CTAAAGGGAG	300
	CCCCCGATTT AGAGCTTGAC GGGGAAAGCC GGCGAACGTG GCGAGAAAGG AAGGAAAGAA	360
15	AGCGAAAGGA GCGGGCGCTA GGGCGCTGGC AAGTGTAGCG GTCAACGCTGC GCGTAACCAC	420
	CACACCCGCC GCGCTTAATG CGCCGCTACA GGGCGCGTCG CGCCATTGCG CATTCAAGGCT	480
20	GCGCAACTGT TGGGAAGGGC GATCGGTGCCG GGCTCTTCG CTATTACGCC AGCTGGCGAA	540
	AGGGGGATGT GCTGCAAGGC GATTAAGTTG GGTAACGCCA GGTTTTCCC AGTCACGACG	600
	TTGTAAAACG ACGGCCAGTG AATTGTAATA CGACTCACTA TAGGGCGAAT TGGGTACCGG	660
25	GCCCCCCCCTC GAGGTCGACG GTATCGATAA GCTTGATGAT CCTTATTGTT ATAGTTCATC	720
	CATGCCATGT GTAATCCCAG CAGCTGTTAC AAACCTCAAGA AGGACCATGT GGTCTCTCTT	780
30	TTCGTTGGGA TCTTTCGAAA GGGCAGATTG TGTTGGACAGG TAATGGTTGT CTGGTAAAAG	840
	GACAGGGCCA TCGCCAATTG GAGTATTTTG TTGATAATGG TCTGCTAGTT GAACGCTTCC	900
	ATCTTCAATG TTGTGTCTAA TTTTGAAGTT AACTTTGATT CCATTCTTTT GTTTGTCTGC	960
35	CATGATGTAT ACATTGTGTG AGTTATAGTT GTATTCCAAT TTGTGTCCAA GAATGTTCC	1020
	ATCTTCTTTA AAATCAATACT CTTTTAACTC GATTCTATTA ACAAGGGTAT CACCTTCAAA	1080
40	CTTGACTTCA GCACGTGTCT TGTTAGTCCC GTCATCTTG AAAAATATAG TTCTTCTTG	1140
	TACATAACCT TCGGGCATGG CACTCTTGAA AAAGTCATGC CGTTTCATAT GATCCGGTA	1200
	TCTTGAAAAG CATTGAACAC CATAAGAGAA AGTAGTGACA AGTGTGGCC ATGGAACAGG	1260
45	TAGTTTCCA GTAGTGCAAA TAAATTAAG GGTAAGTTT CCGTATGTTG CATCACCTTC	1320
	ACCCCTCTCCA CTGACAGAAA ATTTGTGCC ATTAAACATCA CCATCTAATT CAACAAGAAT	1380
	TGGGACAACCT CCAGTGAAGA GTTCTCTCC TTTGCTAGCC ATTTCTTGCG CGATCGAATT	1440
50	CCTGCAGCCC GGGGGATCCA CTAGTTCTAG AGCGGGCGCC ACCGCGGTGG AGCTCCAGCT	1500
	TTTGTCCCT TTAGTGAGGG TTAATTCCGA GCTTGGCGTA ATCATGGTCA TAGCTGTTTC	1560
55	CTGTGTGAAA TTGTTATCCG CTCACAATTC CACACAACAT ACGAGCCGGA AGCATAAAAGT	1620
	GTAAAGCCTG GGGTGCCTAA TGAGTGAGCT AACTCACATT AATTGCGTTG CGCTCACTGC	1680
60	CCGCTTCCA GTCGGGAAAC CTGTCGTGCC AGCTGCATTA ATGAATCGGC CAACGCCGG	1740
	GGAGAGGCCGG TTTGCGTATT GGGCGCTCTT CCGCTTCCCTC GCTCACTGAC TCGCTGCGCT	1800
	CGGTCTTTCG GCTGCGGCCGA GCGGTATCAG CTCACTCAA GGCGGTAAATA CGGTTATCCA	1860
65	CAGAACATCAGG GGATAAACGCA GGAAAGAACAA TGTGAGCAAA AGGCCAGCAA AAGGCCAGGA	1920
	ACCGTAAAAA GGCGCGTTG CTGGCGTTTT TCCATAGGCT CCGCCCCCT GACGAGCATC	1980
	ACAAAAATCG ACGCTCAAGT CAGAGGTGGC GAAACCCGAC AGGACTATAA AGATACCAGG	2040

	CGTTTCCCCC	TGGAAGCTCC	CTCGTGCCT	CTCCTGTTCC	GACCCTGCCG	CTTACCGGAT	2100
	ACCTGTCCGC	CTTTCTCCCT	TCGGGAAGCG	TGGCGCTTTC	TCATAGCTCA	CGCTGTAGGT	2160
5	ATCTCAGTTC	GGTGTAGGTC	GTTCGCTCCA	AGCTGGGCTG	TGTGCACGAA	CCCCCCGTT	2220
	AGCCCGACCG	CTGCGCTTA	TCCGGTAACT	ATCGTCTTGA	GTCCAACCCG	GTAAGACACG	2280
10	ACTTATCGCC	ACTGGCAGCA	GCCACTGGTA	ACAGGATTAG	CAGAGCGAGG	TATGTAGGCG	2340
	GTGCTACAGA	GTTCTTGAAG	TGGTGGCTA	ACTACGGCTA	CACTAGAAGG	ACAGTATTTG	2400
	GTATCTGCGC	TCTGCTGAAG	CCAGTTACCT	TCGGAAAAAG	AGTTGGTAGC	TCTTGATCCG	2460
15	GCAAACAAAC	CACCGCTGGT	AGCGGTGGTT	TTTTTGTGTTG	CAAGCAGCAG	ATTACGCGCA	2520
	GAAAAAAAGG	ATCTCAAGAA	GATCCTTGA	TCTTTTCTAC	GGGGTCTGAC	GCTCAGTGGA	2580
20	ACGAAAACCTC	ACGTTAACGGG	ATTTTGGTCA	TGAGATTATC	AAAAAGGATC	TTCACCTAGA	2640
	TCCTTTAAA	TTAAAAATGA	AGTTTTAAAT	CAATCTAAAG	TATATATGAG	AAAACTTGGT	2700
	CTGACAGTTA	CCAATGCTTA	ATCAGTGAGG	CACCTATCTC	AGCGATCTGT	CTATTCGTT	2760
25	CATCCATAGT	TGCCCTGACTC	CCCCTCGTGT	AGATAACTAC	GATAACGGAG	GGCTTACCAT	2820
	CTGGCCCCAG	TGCTGCAATG	ATACCGCGAG	ACCCACGCTC	ACCGGCTCCA	GATTATCAG	2880
30	CAATAAACCA	GCCAGCCGGA	AGGGCCGAGC	GCAGAAGTGG	TCCTGCAACT	TTATCCGCCT	2940
	CCATCCAGTC	TATTAATTGT	TGCCGGGAAG	CTAGAGTAAG	TAGTCGCCA	GTTAATAGTT	3000
	TGCGCAACGT	TGTTGCCATT	GCTACAGGCA	TCGTGGTGT	ACGCTCGTCG	TTGGTATGG	3060
35	CTTCATTCA	CTCCGGTTCC	CAACGATCAA	GGCGAGTTAC	ATGATCCCCC	ATGTTGTGCA	3120
	AAAAAGCGGT	TAGCTCCTTC	GGTCCTCCGA	TCGTTGTCAG	AAGTAAGTTG	GCCGCAGTGT	3180
40	TATCACTCAT	GGTTATGGCA	GCACTGCATA	ATTCTCTTAC	TGTCATGCCA	TCCGTAAGAT	3240
	GCTTTCTGT	GACTGGTGAG	TACTCAACCA	AGTCATTCTG	AGAATAGTGT	ATGCGCGAC	3300
	CGAGTTGCTC	TTGCCCGGCG	TCAATACGGG	ATAATACCCG	GCCACATAGC	AGAACTTTAA	3360
45	AAGTGCTCAT	CATTGGAAAA	CGTTCTTCGG	GGCGAAAAC	CTCAAGGATC	TTACCGCTGT	3420
	TGAGATCCAG	TTCGATGTAA	CCCACTCGTG	CACCCAACTG	ATCTTCAGCA	TCTTTACTT	3480
50	TCACCAGCGT	TTCTGGGTGA	GCAAAACAG	GAAGGAAAA	TGCCGCAGAA	AAGGAAATAA	3540
	GGGCGACACG	GAAATGTTGA	ATACTCATA	TCTTCCTTTT	TCAATATTAT	TGAAGCATT	3600
	ATCAGGGTTA	TTGTCTCATG	AGCGGATACA	TATTGAAATG	TATTTAGAAA	ATAAAACAAA	3660
55	TAGGGGTTCC	GCCGACATTT	CCCCGAAAAG	TGCCACCTG			3699

## (2) INFORMATION FOR SEQ ID NO: 7:

60 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 6361 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

65 (ii) MOLECULE TYPE: DNA

(ix) FEATURE:  
 (A) NAME/KEY: -

(B) LOCATION: 1..6361  
 (D) OTHER INFORMATION: /note= "pFRED13"

5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:	
10	TTCTCATGTT TGACAGCTTA TCATCGATAA GCTTTAACATGC GGTAGTTTAT CACAGTTAAA	60
15	TTGCTAACGC AGTCAGGCAC CGTGTATGAA ATCTAACAAAT GCGCTCATCG TCATCCTCGG	120
20	CACCGTCACC CTGGATGCTG TAGGCATAGG CTTGGTTATG CCGGTACTGC CGGGCCTCTT	180
25	GCGGGATATC CGGATATAGT TCCTCCTTTC AGCAAAAAAC CCCTCAAGAC CCGTTTAGAG	240
30	GCCCCAAGGG GTTATGCTAG TTATTGCTCA GCGGTGGCAG CAGCCAACTC AGCTTCCTTT	300
35	CGGGCTTTGT TAGCAGCCGG ATCCTTATTT GTATAGTTCA TCCATGCCAT GTGTAATCCC	360
40	AGCAGCTGTT ACAAAACTCAA GAAGGACCAT GTGGTCTCTC TTTTCGTTGG GATCTTCGA	420
45	AAGGGCAGAT TGTGTGGACA GGTAATGGTT GTCTGGTAAA AGGACAGGGC CATGCCAAT	480
50	TGGAGTATTT TGTTGATAAT GGTCTGCTAG TTGAACGCTT CCATCTTCAA TGTTGTGTCT	540
55	AATTTTGAAG TTAACCTTGAA TTCCATTCTT TTGTTTGTCT GCCATGATGT ATACATTGTG	600
60	TGAGTTATAG TTGTATTCCA ATTTGTGTCC AAGAAATGTTT CCATCTTCTT TAAAATCAAT	660
65	ACCTTTAAC TCGATTCTAT TAACAAGGGT ATCACCTTCA AACITGACTT CAGCACGTGT	720
70	CTTGTAGTTC CCGTCATCTT TGAAAATAT AGTTCTTCC TGTACATAAC CTTGGGCAT	780
75	GGCACTCTTG AAAAAGTCAT GCCGTTTCAT ATGATCCGGG TATCTTGGAA AGCATTGAAC	840
80	ACCATAAGAG AAAGTAGTGA CAAGTGTGG CCATGGAACA GGTAGTTTC CAGTAGTGCA	900
85	AATAAAATTAA AGGGTAAGTT TTCCGTATGT TGCATCACCT TCACCCCTCTC CACTGACAGA	960
90	AAATTTGTGC CCATTAACAT CACCATCTAA TTCAACAAGA ATTGGGACAA CTCCAGTGAA	1020
95	GAGTTCTTCT CCTTTGCTAG CCATATGTAT ATCTCCTTCT TAAAGTTAAA CAAAATTATT	1080
100	TCTAGAGGGG AATTGTTATC CGCTCACAAAT TCCCCTATAG TGAGTCGTAT TAATTGCGC	1140
105	GGATCGAGAT CTCGATCCTC TACGCCGGAC GCATCGTGGC CGGCATCACCC GGCGCCACAG	1200
110	GTGCGGTTGC TGGCGCTAT ATCGCCGACA TCACCGATGG GGAAGATCGG GCTCGCCACT	1260
115	TCGGGCTCAT GAGCGCTTGT TTCCGGTGG GTATGGTGGC AGGCCCCGTG GCCGGGGAC	1320
120	TGTTGGCGC CATCTCCTTG CATGCACCAT TCCTTGCCTGC GGCGGTGCTC AACGGCCTCA	1380
125	ACCTACTACT GGGCTGCTTC CTAATGCAGG AGTGGATAA GGGAGAGCGT CGAGATCCCG	1440
130	GACACCATCG AATGGCGAA AACCTTCGC GGTATGGCAT GATAGCGCCC GGAAGAGAGT	1500
135	CAATTCAAGGG TGGTGAATGT GAAACCAGTA ACCTTATACG ATGTCGCAGA GTATGCCGT	1560
140	GTCTCTTATC AGACCGTTTC CGCGTGGTG AACCAAGGCCA GCCACGTTTC TGCAGAAACG	1620
145	CGGGAAAAAG TGGAAGCGGC GATGGCGGAG CTGAATTACA TTCCCAACCG CGTGGCACAA	1680
150	CAACTGGCGG GCAAACAGTC GTTGTGATT GGCGTTGCCA CCTCCAGTCT GGCCCTGCAC	1740
155	GCGCCGTCGC AAATTGTGCGC GGCGATTAAA TCTCGCGCCG ATCAACTGGG TGCCAGCGTG	1800
160	GTGGTGTGCA TGGTGAACG AAGCGGCGTC GAAGCCTGTA AAGCGGCGGT GCACAATCTT	1860
165	CTCGCGCAAC GCGTCAGTGG GCTGATCATT AACTATCCGC TGGATGACCA GGATGCCATT	1920

	GCTGTGGAAG CTGCCTGCAC TAATGTTCCG GCGTTATTTC TTGATGTCTC TGACCAGACA	1980
	CCCATCAACA GTATTATTTT CTCCCAGAA GACGGTACGC GACTGGCGT GGAGCATCTG	2040
5	GTCGCATTGG GTCACCAGCA AATCGCGCTG TTAGCGGCC CATTAAAGTC TGTCTCGCG	2100
	CGTCTGCGTC TGGCTGGCTG GCATAAATAT CTCACTCGCA ATCAAATTCA GCCGATAGCG	2160
10	GAACGGGAAG GCGACTGGAG TGCCATGTCC GGTGTTCAAC AAACCATGCA AATGCTGAAT	2220
	GAGGGCCTCG TTCCCCACTGC GATGCTGGTT GCCAACGATC AGATGGCGT GGGCGCAATG	2280
	CGCGCCATTA CCGAGTCCGG GCTGCGCGT GGTGCGGATA TCTCGGTAGT GGGATACGAC	2340
15	GATACCGAAG ACAGCTCATG TTATATCCCG CCGTTAACCA CCATCAAACA GGATTTCGC	2400
	CTGCTGGGGC AAACCAGCGT GGACCCCTTG CTGCAACTCT CTCAGGGCCA GGCGGTGAAG	2460
20	GGCAATCAGC TGTTGCCCCG CTCACTGGTG AAAAGAAAAA CCACCCCTGGC GCCCAATACG	2520
	CAAACCGCCT CTCCCCCGCG CTTGGCCGAT TCATTAATGC AGCTGGCACG ACAGGTTCC	2580
	CGACTGGAAA CGGGGCAGTG AGCGAACGC ATTAAATGTA AGTTAGCTCA CTCATTAGGC	2640
25	ACCGGGATCT CGACCGATGC CCTTGAGAGC CTTAACCCCA GTCAGCTCCT TCCGGTGGGC	2700
	GCAGGGCATG ACTATCGTCG CCGCACTTAT GACTGTCTTC TTTATCATGC AACTCGTAGG	2760
30	ACAGGTGCCG GCAGCGCTCT GGGTCATTTT CGCGGAGGAC CGCTTCGCT GGAGCGCGAC	2820
	GATGATCGGC CTGTCGCTTG CGGTATTCGG AATCTTGAC GCCCTCGCTC AAGCCTTCGT	2880
	CACTGGTCCC GCCACCAAAC GTTTCGGCGA GAAGCAGGCC ATTATCGCCG GCATGGCGGC	2940
35	CGACCGCCTG GGCTACGTCT TGCTGGCGTT CGCGACGCGA GGCTGGATGG CCTTCCCCAT	3000
	TATGATTCTT CTCGCTTCCG GCGGCATCGG GATGCCCGCG TTGCAGGCCA TGCTGTCCAG	3060
40	GCAGGTAGAT GACGACCATC AGGGACAGCT TCAAGGATCG CTCGCGCTC TTACCAAGCCT	3120
	AACTTCGATC ACTGGACCGC TGATCGTCAC GGCGATTTAT GCCGCCTCGG CGAGCACATG	3180
	GAACGGGTTG GCATGGATTG TAGGCGCCGC CCTATACCTT GTCTGCCTCC CCGCGTTGGC	3240
45	TCGCGGTGCA TGGAGCCGGG CCACCTCGAC CTGAATGGAA GCCGGCGCA CCTCGCTAAC	3300
	GGATTCACCA CTCCAAGAAT TGGAGCCAAT CAATTCTTGC GGAGAACTGT GAATGCCAA	3360
50	ACCAACCCCTT GGCAGAACAT ATCCATCGCG TCCGCCATCT CCAGCAGCCG CACCGGGCGC	3420
	ATCTCGGGCA GCGTTGGGTC CTGGCCACGG GTGCGCATGA TCGTGCCTCT GTCGTTGAGG	3480
	ACCCGGCTAG GCTGGGGGGG TTGGCTTACT GGTTAGCAGA ATGAATCACC GATAACGGAG	3540
55	CGAACGTGAA GCGACTGCTG CTGAAAACG TCTGCGACCT GAGCAACAAAC ATGAATGGTC	3600
	TTCGGTTTCC GTGTTTCGTA AAGTCTGGAA ACGGGAAAGT CAGGCCCTG CACCAATTATG	3660
60	TTCCGGATCT GCATCGCAGG ATGCTGCTGG CTACCCGTG GAACACCTAC ATCTGTATTA	3720
	ACGAAGCGCT GGCATTGACC CTGAGTGATT TTTCTCTGGT CCCGCCGCAT CCATACCGCC	3780
	AGTTGTTTAC CCTCACAAACG TTCCAGTAAC CGGGCATGTT CATCATCAGT AACCCGTATC	3840
65	GTGAGCATCC TCTCTCGTTT CATCGGTATC ATTACCCCA TGAACAGAAA TCCCCCTTAC	3900
	ACGGAGGCAT CAGTGACCAA ACAGGAAAAA ACCGCCCTTA ACATGGCCCG CTTTATCAGA	3960
	AGCCAGACAT TAACGCTTCT GGAGAAACTC AACGAGCTGG ACGGCGATGA ACAGGCAGAC	4020

	ATCTGTGAAT CGCTTCACGA CCACGCTGAT GAGCTTTACC GCAGCTGCCT CGCGCGTTTC	4080
	GGTGATGACG GTGAAAACCT CTGACACATG CAGCTCCCG AGACGGTCAC AGCTTGTCTG	4140
5	TAAGCGGATG CCGGGAGCAG ACAAGCCCAGT CAGGGCGCGT CAGCGGGTGT TGGCGGGTGT	4200
	CGGGCGCGAG CCATGACCCA GTCACCGTAGC GATAGCGGAG TGTATACTGG CTTAACTATG	4260
10	CGGCATCAGA GCAGATTGTA CTGAGAGTGC ACCATATATG CGGTGTGAAA TACCGCACAG	4320
	ATGCGTAAGG AGAAAATACC GCATCAGGCG CTCTTCCGCT TCCTCGCTCA CTGACTCGCT	4380
	GCGCTCGGTC GTTCGGCTGC GGCGAGCGGT ATCAGCTCAC TCAAAGGCGG TAATACGGTT	4440
15	ATCCACAGAA TCAGGGGATA ACGCAGGAAA GAACATGTGA GCAAAGGCC AGCAAAGGC	4500
	CAGGAACCGT AAAAAGGCCG CGTTGCTGGC GTTTTCCAT AGGCTCCGCC CCCCTGACGA	4560
20	GCATCACAAA AATCGACGCT CAAGTCAGAG GTGGCGAAC CCGACAGGAC TATAAAGATA	4620
	CCAGGCCTTT CCCCCCTGGAA GCTCCCTCGT GCGCTCTCCT GTTCCGACCC TGCCGCTTAC	4680
	CGGATACCTG TCCGCCCTTC TCCCTTCGGG AAGCGTGGCG CTTTCTCATA GCTCACGCTG	4740
25	TAGGTATCTC AGTCGGTGT AGGTCGTTCG CTCCAAGCTG GGCTGTGTGC ACGAACCCCC	4800
	CGITCAGCCC GACCGCTGCG CCTTATCCGG TAACTATCGT CTTGAGTCCA ACCCGGTAAG	4860
30	ACACGACTTA TCGCCACTGG CAGCAGCCAC TGGTAACAGG ATTAGCAGAG CGAGGTATGT	4920
	AGGCGGTGCT ACAGAGTTCT TGAAGTGGTG GCCTAACTAC GGCTACACTA GAAGGACAGT	4980
	ATTTGGTATC TGCGCTCTGC TGAAGCCAGT TACCTTCGGA AAAAGAGTTG GTAGCTCTTG	5040
35	ATCCGGCAAA CAAACCACCG CTGGTAGCGG TGGTTTTTTT GTTTGCAAGC AGCAGATTAC	5100
	GCGCAGAAAA AAAGGATCTC AAGAAGATCC TTGATCTTT TCTACGGGGT CTGACGCTCA	5160
	GTGGAACGAA AACTCACGTT AAGGGATTTT GGTACATGAGA TTATCAAAA GGATCTTCAC	5220
40	CTAGATCCTT TAAATTAAA AATGAAGTTT TAAATCAATC TAAAGTATAT ATGAGTAAAC	5280
	TTGGTCTGAC AGTTACCAAT GCTTAATCAG TGAGGCACCT ATCTCAGCGA TCTGCTTATT	5340
45	TCGTTCATCC ATAGTTGCCT GACTCCCCGT CGTGTAGATA ACTACGATAAC GGGAGGGCTT	5400
	ACCATCTGGC CCCAGTGCTG CAATGATAACC GCGAGACCCA CGCTCACCCG CTCCAGATT	5460
50	ATCAGCAATA AACCAAGCCAG CGGAAGGGC CGAGCGCAGA AGTGGTCCTG CAACTTTATC	5520
	CGCCTCCATC CAGTCTATTA ATTGTTGCCG GGAAGCTAGA GTAAGTAGTT CGCCAGTTAA	5580
	TAGTTGCGC AACGTTGTTG CCATTGCTGC AGGCATCGTG GTGTACCGCT CGTCGTTGG	5640
55	TATGGCTTCA TTCAGCTCCG GTTCCCAACG ATCAAGGCGA GTTACATGAT CCCCCATGTT	5700
	GTGCAAAAAA GCGGTTAGCT CCTTCGGTCC TCCGATCGTT GTCAGAAAGTA AGTGGCCGC	5760
60	AGTGTATCA CTCATGGTTA TGGCAGCACT GCATAATTCT CTTACTGTCA TGCCATCCGT	5820
	AAGATGCTTT TCTGTGACTG GTGAGTACTC AACCAAGTCA TTCTGAGAAT AGTGTATGCG	5880
	GCGACCGAGT TGCTCTTGGCC CGGCGTCAAC ACGGGATAAT ACCGCGCCAC ATAGCAGAAC	5940
65	TTTAAAAGTG CTCATCATTG GAAAACGTTT TTCGGGGCGA AAACTCTCAA GGATCTTACC	6000
	GCTGTTGAGA TCCAGTTCGA TGTAACCCAC TCGTGCACCC AACTGATCTT CAGCATTCTT	6060
	TACTTTCACC AGCGTTCTG GGTGAGCAA AACAGGAAGG CAAAATGCCG CAAAAAAGGG	6120

	AATAAGGGCG ACACGGAAAT GTTGAATACT CATACTCTTC CTTTTCAAT ATTATTGAAG	6180
	CATTATCAG GGTTATTGTC TCATGAGCGG ATACATATTT GAATGTATTT AGAAAATAA	6240
5	ACAAATAGGG GTTCCGGCGA CATTCCCCG AAAAGTGCCA CCTGACGTCT AAGAAACCAT	6300
	TATTATCATG ACATTAACCT ATAAAAATAG GCGTATCACG AGGCCCTTTC GTCTTCAAGA	6360
	A	6361
10		

## (2) INFORMATION FOR SEQ ID NO:8:

15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 48 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
20	(ii) MOLECULE TYPE: DNA	
25	(ix) FEATURE: (A) NAME/KEY: - (B) LOCATION: 1..48 (D) OTHER INFORMATION: /note= "oligonucleotide #17422"	
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8: CAATTTGTGT CCCAGAATGT TGCCATCTTC CTTGAAGTCA ATACCTTT	48
35	(2) INFORMATION FOR SEQ ID NO:9:	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 47 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
45	(ii) MOLECULE TYPE: DNA	

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..47
- (D) OTHER INFORMATION: /note= "oligonucleotide #17423"

5

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

GTCTTGTAGT TGCCGTCATC TTTGAAGAAG ATGCTCCTTT CCTGTAC

47

10

## (2) INFORMATION FOR SEQ ID NO:10:

15 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 52 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

20

## (ii) MOLECULE TYPE: DNA

## (ix) FEATURE:

25 (A) NAME/KEY: -  
(B) LOCATION: 1..52  
(D) OTHER INFORMATION: /note= "oligonucleotide #17424"

30

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

CATGGAACAG GCAGTTGCC AGTAGTGCAG ATGAACTTCA GGGTAAGTTT TC

52

35

## (2) INFORMATION FOR SEQ ID NO:11:

40 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 40 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA

45

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..40
- (D) OTHER INFORMATION: /note= "oligonucleotide #17425"

50

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

CTCCACTGAC AGAGAACTTG TGGCCGTTAA CATCACCATC

40

55

## (2) INFORMATION FOR SEQ ID NO:12:

60 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 47 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

65

## (ii) MOLECULE TYPE: DNA

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..47
- (D) OTHER INFORMATION: /note= "oligonucleotide #17426"

5

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

CCATCTCAA TGTTGTGGCG GGTCTTGAAG TTCACTTTGA TTCCATT  
10

47

## (2) INFORMATION FOR SEQ ID NO:13:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 41 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

20

## (ii) MOLECULE TYPE: DNA

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..41
- (D) OTHER INFORMATION: /note= "oligonucleotide #17465"

30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:  
CGATAAGCTT GAGGATCCTC AGTTGTACAG TTCAATCCATG C

41

## (2) INFORMATION FOR SEQ ID NO:14:

35

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 849 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

40

## (ii) MOLECULE TYPE: DNA

45

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..849
- (D) OTHER INFORMATION: /note= "pBSGFPsg11"

50

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

ATGACCATGA TTACGCCAAG CTCGGAATTA ACCCTCACTA AAGGGAACAA AAGCTGGAGC	60
TCCACCGCGG TGGCGGCCGC TCTAGAACTA GTGGATCCCC CGGGCTGCAG GAATTCGATC	120
GCGCAAGAAA TGGCTAGCAA AGGAGAAGAA CTCTTCACTG GAGTTGTCCC AATTCTTGTT	180
GAATTAGATG GTGATGTTAA CGGCCACAAG TTCTCTGTCA GTGGAGAGGG TGAAGGTGAT	240
GCAACATACG GAAAACTTAC CCTGAAGTTC ATCTGCACTA CTGGCAAACCT GCCTGTTCCA	300
TGGCCAAACAC TTGTCACTAC TCTCTCTTAT GGTGTTCAAT GCTTTTCAAG ATACCCGGAT	360
CATATGAAAC GGCATGACTT TTTCAAGAGT GCCATGCCCG AAGGTTATGT ACAGGAAAGG	420
ACCATCTTCT TCAAAGATGA CGGCAACTAC AAGACACGTG CTGAAGTCAA GTTTGAAGGT	480
GATAACCCTTG TTAATAGAAT CGAGTTAAAA GGTATTGACT TCAAGGAAGA TGGCAACATT	540

	CTGGGACACA AATTGGAATA CAACTATAAC TCACACAATG TATACATCAT GGCAGACAAA	600
	CAAAAGAATG GAATCAAAGT GAACTTCAG ACCCGCCACA ACATTGAAGA TGGAAGCGTT	660
5	CAACTAGCAG ACCATTATCA ACAAAATACT CCAATTGGCG ATGGCCCTGT CCTTTTACCA	720
	GACAACCATT ACCTGTCCAC ACAATCTGCC CTTCGAAAG ATCCCAACGA AAAGAGAGAC	780
10	CACATGGTCC TTCTTGAGTT TGTAACAGCT GCTGGGATTACACATGGCAT GGATGAAC	840
	TACAACGTGA	849

15 (2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 720 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

25 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..720  
 (D) OTHER INFORMATION: /note= "SG12"

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

	ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
35	GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
	GGAAAAACTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCTGTTC ATGGCCAACA	180
40	CTTGTCACTA CTCTCTCTTA TGGTGTCAA TGCTTTCAA GATACCCGGA TCATATGAAA	240
	CGGCATGACT TTTTCAAGAG TGCCATGCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
	TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAAGG TGATACCC	360
45	GTAAATAGAA TCGAGTTAAA AGGTATTGAT TTTAAAGAAG ATGGAAACAT TCTTGGACAC	420
	AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
50	GGAAATCAAAG TTAACCTCAA AATTAGACAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
	GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTACCA AGACAACCAT	600
	TACCTGTCCA CACAATCTGC CCTTTCGAAA GATCCCACG AAAAGAGAGA CCACATGGTC	660
55	CTTCTTGAGT TTGTAACAGC TGCTGGGATT ACACATGGCA TGGATGAAC	720
	ATACAAATAA	

(2) INFORMATION FOR SEQ ID NO:16:

60 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 720 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

65 (ii) MOLECULE TYPE: DNA

(ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..720  
 (D) OTHER INFORMATION: /note= "SG11"

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

	ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
10	GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
	GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCGTGTCC ATGCCAACAA	180
15	CTTGTCACTA CTCTCTCTTA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
	CGGCATGACT TTTTCAAGAG TGCCATGCCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
	TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTGAAGG TGATACCCCTT	360
20	GTAAATAGAA TCGAGTTAAA AGGTATTGAC TTCAAGGAAG ATGGCAACAT TCTGGGACAC	420
	AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
25	GGAATCAAAG TGAACATTCAA GACCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
	GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCCTTTACC AGACAACCCT	600
	TACCTGTCCA CACAATCTGC CCTTTCGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
30	CTTCTTGAGT TTGTAACAGC TGCTGGGATT ACACATGGCA TGGATGAACT GTACAACGTGA	720

(2) INFORMATION FOR SEQ ID NO:17:

35

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 720 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

40

(ii) MOLECULE TYPE: DNA

45

(ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..720  
 (D) OTHER INFORMATION: /note= "SG25"

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

	ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
55	GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
	GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCGTGTCC ATGCCAACAA	180
	CTAGTCACTA CTCTGTGCTA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
60	CGGCATGACT TTTTCAAGAG TGCCATGCCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
	TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTGAAGG TGATACCCCTT	360
65	GTAAATAGAA TCGAGTTAAA AGGTATTGAC TTCAAGGAAG ATGGCAACAT TCTGGGACAC	420
	AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
	GGAATCAAAG TGAACATTCAA GACCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540

GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTACC AGACAACCAT	600
TACCTGTCCA CACAATCTGC CCTTCGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
5 CTTCTTGAGT TTGTAACAGC TGCTGGGATT ACACATGGCA TGGATGAAC GTACAACGTGA	720

## (2) INFORMATION FOR SEQ ID NO:18:

10 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 40 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

15 (ii) MOLECULE TYPE: DNA

20 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..40  
 (D) OTHER INFORMATION: /note= "oligonucleotide #18217"

## 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

CATTGAACAC CATAGCACAG AGTAGTGACT AGTGTGGCC	40
--------------------------------------------	----

## 30 (2) INFORMATION FOR SEQ ID NO:19:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 720 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

40 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..720  
 (D) OTHER INFORMATION: /note= "SB42"

## 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

50 ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCTGTTCATGGCCAACAA	180
55 CTAGTCACTA CTCTCTCTCA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
CGGCATGACT TTTTCAAGAG TGCCATGCCA GAAGGTTATG TACAGGAAAG GACCATCTTC	300
60 TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAGG TGATAACCTT	360
GTAAATAGAA TCGAGTTAAA AGGTATTGAT TTTAAAGAAG ATGGAAACAT TCTTGGACAC	420
AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
65 GGAATCAAAG TTAACCTCAA ATTAGACAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTACC AGACAACCAT	600
TACCTGTCCA CACAATCTGC CCTTCGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660

CTTCTTGAGT TTGTAACAGC TGCTGGATT ACACATGGCA TGGATGAAC ATACAAATAA 720

5 (2) INFORMATION FOR SEQ ID NO:20:

10 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 40 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

15 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..40  
 (D) OTHER INFORMATION: /note= "oligonucleotide #bio25"

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

CATTGAACAC CATGAGAGAG AGTAGTGACT AGTGTTGGCC 40

25 (2) INFORMATION FOR SEQ ID NO:21:

30 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 720 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

35 (ii) MOLECULE TYPE: DNA

40 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..720  
 (D) OTHER INFORMATION: /note= "SB49"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

45	ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
	GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
50	GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAC TGCCTGTTCC ATGGCCAACA	180
	CTAGTCACTA CTTTCTCTCA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
	CGGCATGACT TTTTCAAGAG TGCCATGCCA GAAGGTTATG TACAGGAAAG GACCATCTTC	300
55	TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAAGG TGATACCCCTT	360
	GTAAATAGAA TCGAGTTAAA AGGTATTGAT TTTAAAGAAG ATGGAAACAT TCTTGGACAC	420
60	AAATTGGAAT ACAACTATAA CTCACACAAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
	GGAATCAAAG CGAACTTCAA GATCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
	GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTACCA AGACAACCAT	600
65	TACCTGTCCA CACAATCTGC CCTTTGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
	CTTCTTGAGT TTGTAACAGC TGCTGGATT ACACATGGCA TGGATGAAC ATACAAATAA	720

## (2) INFORMATION FOR SEQ ID NO:22:

5                   (i) SEQUENCE CHARACTERISTICS:  
                      (A) LENGTH: 44 base pairs  
                      (B) TYPE: nucleic acid  
                      (C) STRANDEDNESS: single  
                      (D) TOPOLOGY: linear

10                  (ii) MOLECULE TYPE: DNA

15                  (ix) FEATURE:  
                      (A) NAME/KEY: -  
                      (B) LOCATION: 1..44  
                      (D) OTHER INFORMATION: /note= "oligonucleotide #19059"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

20                  CTTCAATGTT GTGGCGGATC TTGAAGTCG CTTTGATTCC ATTC

44

## (2) INFORMATION FOR SEQ ID NO:23:

25                  (i) SEQUENCE CHARACTERISTICS:  
                      (A) LENGTH: 40 base pairs  
                      (B) TYPE: nucleic acid  
                      (C) STRANDEDNESS: single  
                      (D) TOPOLOGY: linear

30                  (ii) MOLECULE TYPE: DNA

35                  (ix) FEATURE:  
                      (A) NAME/KEY: -  
                      (B) LOCATION: 1..40  
                      (D) OTHER INFORMATION: /note= "oligonucleotide #bio24."

40                  (xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

CATTGAACAC CATGAGAGAA AGTAGTGACT AGTGTGGCC

40

## 45                  (2) INFORMATION FOR SEQ ID NO:24:

50                  (i) SEQUENCE CHARACTERISTICS:  
                      (A) LENGTH: 720 base pairs  
                      (B) TYPE: nucleic acid  
                      (C) STRANDEDNESS: single  
                      (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

55

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..720
- (D) OTHER INFORMATION: /note= "SB50"

5

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCCTGTTCC ATGGCCAACA	180
CTAGTCACTA CTCTCTCTCA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
CGGCATGACT TTTTCAAGAG TGCCATGCCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAAGG TGATACCCCTT	360
GTAAATAGAA TCGAGTTAAA AGGTATTGAT TTTAAAGAAG ATGGAAACAT TCTGGACAC	420
AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
GGAATCAAAG CGAACATTCAA GATCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
GACCATTATC AACAAAATAC TCCAATTGGC GATGCCCTG TCCTTTTACC AGACAACCAT	600
TACCTGTCCA CACAATCTGC CCTTTCGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
CTTCTTGAGT TTGTAACAGC TGCTGGGATT ACACATGGCA TGGATGAAC ATACAAATAA	720

35

## (2) INFORMATION FOR SEQ ID NO:25:

40

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1521 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA

45

## (ix) FEATURE:

- (A) NAME/KEY: -
- (B) LOCATION: 1..1521
- (D) OTHER INFORMATION: /note= "pCMVgfol1"

50

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCCTGTTCC ATGGCCAACA	180
CTTGTCACTA CTCTCTCTTA TGGTGTCAA TGCTTTCAA GATAACCGGA TCATATGAAA	240
CGGCATGACT TTTTCAAGAG TGCCATGCCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAAGG TGATACCCCTT	360
GTAAATAGAA TCGAGTTAAA AGGTATTGAC TTCAAGGAAG ATGGAAACAT TCTGGACAC	420
AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAT	480
GGAATCAAAG TGAACATTCAA GACCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540

	GACCATTATC AACAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTACCG AGACAACCAT	600
	TACCTGTCCA CACAATCTGC CCTTCGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
5	CTTCTTGAGT TTGTAAACAGC TGCTGGGATT ACACATGGCA TGGATGAACCT GTACAACGGT	720
	GCTGGTGCTA TCGAACAAAGA TGGATTGCAC CCAGGGTCTC CGGCCGCTTG GGTGGAGAGG	780
10	CTATTGGCT ATGACTGGGC ACAACAGACA ATCGGCTGCT CTGATGCCGC CGTGTTCGG	840
	CTGTCAGCGC AGGGGCGCCC GGTTCTTTT GTCAAGACCG ACCTGTCCGG TGCCCTGAAT	900
	GAAC TGCAAGG CAGG GCGCTATCG TGGCTGGCCA CGACGGCGT TCCTTGCGCA	960
15	GCTGTGCTCG ACGTTGTAC TGAAGCGGGA AGGGACTGGC TGCTATTGGG CGAAGTGCCG	1020
	GGGCAGGATC TCCTGTACATC TCACCTTGCT CCTGCCGAGA AAGTATCCAT CATGGCTGAT	1080
20	GCAATGCGGC GGCTGCATAAC GCTTGATCCG GCTACCTGCC CATTGACCA CCAAGCGAAA	1140
	CATCGCATCG AGCGAGCACG TACTCGGATG GAAGCCGGTC TTGTCGATCA GGATGATCTG	1200
	GACGAAGAGC ATCAGGGGCT CGCGCCAGCC GAACTGTTCG CCAGGCTCAA GGCGCGCATG	1260
25	CCCGACGGCG AGGATCTCGT CGTGACCCAT GGCATGCCCT GCTTGCCGAA TATCATGGTG	1320
	GAAAATGGCC GCTTTTCTGG ATTCAATCGAC TGTGGCCGGC TGGGTGTGGC GGACCGCTAT	1380
30	CAGGACATAG CGTTGGCTAC CCGTGATATT GCTGAAGAGC TTGGCGGCGA ATGGGCTGAC	1440
	CGCTTCCTCG TGCTTTACGG TATCGCCGCT CCCGATTGCG AGCGCATCGC CTTCTATCGC	1500
	CTTCTTGACG AGTTCTTCTG A	1521

35

## (2) INFORMATION FOR SEQ ID NO:26:

40 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 4 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS:  
 (D) TOPOLOGY: linear

45 (ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

50 Gly Ala Gly Ala  
1

## (2) INFORMATION FOR SEQ ID NO:27:

55 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 32 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

60 (ii) MOLECULE TYPE: DNA

5 (ix) FEATURE:  
      (A) NAME/KEY: -  
      (B) LOCATION: 1..32  
      (D) OTHER INFORMATION: /note= "primer Bio51"

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:  
CGCGGATCCT TCGAACAAAGA TGGATTGCAC GC

15 (2) INFORMATION FOR SEQ ID NO:28:  
      (i) SEQUENCE CHARACTERISTICS:  
          (A) LENGTH: 34 base pairs  
          (B) TYPE: nucleic acid  
          (C) STRANDEDNESS: single  
          (D) TOPOLOGY: linear

20 (ii) MOLECULE TYPE: DNA

25 (ix) FEATURE:  
      (A) NAME/KEY: -  
      (B) LOCATION: 1..34  
      (D) OTHER INFORMATION: /note= "primer Bio52"

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:  
CCGGAATTCT CAGAAGAACT CGTCAAGAAG GCGA

35 (2) INFORMATION FOR SEQ ID NO:29:  
      (i) SEQUENCE CHARACTERISTICS:  
          (A) LENGTH: 46 base pairs  
          (B) TYPE: nucleic acid  
          (C) STRANDEDNESS: single  
          (D) TOPOLOGY: linear

40 (ii) MOLECULE TYPE: DNA

45 (ix) FEATURE:  
      (A) NAME/KEY: -  
      (B) LOCATION: 1..46  
      (D) OTHER INFORMATION: /note= "primer Bio49"

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:  
GGCGCGCAAG AAATGGCTAG CAAAGGAGAA GAACTCTTCA CTGGAG

55 (2) INFORMATION FOR SEQ ID NO:30:  
      (i) SEQUENCE CHARACTERISTICS:  
          (A) LENGTH: 46 base pairs  
          (B) TYPE: nucleic acid  
          (C) STRANDEDNESS: single  
          (D) TOPOLOGY: linear

60 (ii) MOLECULE TYPE: DNA

## (ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..46  
 (D) OTHER INFORMATION: /note= "primer Bio50"

5

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

10 CCCATCGATA GCACCAGCAC CGTTGTACAG TTCAATCCATG CCATGT

46

## (2) INFORMATION FOR SEQ ID NO:31:

15 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 1521 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

20 (ii) MOLECULE TYPE: DNA

## (ix) FEATURE:

25 (A) NAME/KEY: -  
 (B) LOCATION: 1..1521  
 (D) OTHER INFORMATION: /note= "pPGKgfo25"

## 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

ATGGCTAGCA AAGGAGAAGA ACTCTTCACT GGAGTTGTCC CAATTCTTGT TGAATTAGAT	60
GGTGATGTTA ACGGCCACAA GTTCTCTGTC AGTGGAGAGG GTGAAGGTGA TGCAACATAC	120
35 GGAAAACCTTA CCCTGAAGTT CATCTGCACT ACTGGCAAAC TGCCCTGTTCC ATGGCCAACA	180
CTAGTCACTA CTCTGTGCTA TGGTGTCAA TGCTTTCAA GATACCCGGA TCATATGAAA	240
40 CGGCATGACT TTTTCAAGAG TGCCATGCCC GAAGGTTATG TACAGGAAAG GACCATCTTC	300
TTCAAAGATG ACGGCAACTA CAAGACACGT GCTGAAGTCA AGTTTGAAGG TGATACCCCTT	360
GTTAATAGAA TCGAGTTAAA AGGTATTGAC TTCAAGGAAG ATGGCAACAT TCTGGGACAC	420
45 AAATTGGAAT ACAACTATAA CTCACACAAT GTATACATCA TGGCAGACAA ACAAAAGAAC	480
GGAATCAAAG TGAACATTCAA GACCCGCCAC AACATTGAAG ATGGAAGCGT TCAACTAGCA	540
50 GACCATTATC ACAAAAATAC TCCAATTGGC GATGGCCCTG TCCTTTTAC AGACAACCAT	600
TACCTGTCCA CACAATCTGC CCTTTGAAA GATCCCAACG AAAAGAGAGA CCACATGGTC	660
55 CTTCTTGAGT TTGTAACAGC TGCTGGGATT ACACATGGCA TGGATGAACGT GTACAACGGT	720
GCTGGTGCTA TCGAACAAAGA TGGATTGCAC GCAGGTTCTC CGGCCGTTG GGTGGAGAGG	780
CTATTGGCT ATGACTGGGC ACAACAGACA ATCGGCTGCT CTGATGCCGC CGTGTCCGG	840
60 CTGTCAGCGC AGGGGCGCCC GGTTCTTTT GTCAAGACCG ACCTGTCCGG TGCCCTGAAT	900
GAAGTGCAGG ACGAGGCAGC GCGGCTATCG TGGCTGGCCA CGACGGGCGT TCCTTGCAC	960
GCTGTGCTCG ACGTTGTAC TGAAGGGGA AGGGACTGGC TGCTATTGGG CGAAGTGCAG	1020
65 GGGCAGGATC TCCTGTCTAC TCACCTTGCT CCTGCCGAGA AAGTATCCAT CATGGCTGAT	1080
GCAATGCGGC GGCTGCATAC GCTTGATCCG GCTACCTGCC CATTGACCA CCAAGCGAAA	1140
CATCGCATCG AGCGAGCACG TACTCGGATG GAAGCCGGTC TTGTCGATCA GGATGATCTG	1200

	GACGAAGAGC ATCAGGGGCT CGCGCCAGCC GAACTGTTCG CCAGGCTCAA GGCGCGCATG	1260
	CCCGACGGCG AGGATCTCGT CGTGACCCAT GGCGATGCCT GCTTGCCGAA TATCATGGTG	1320
5	GAAAATGGCC GCTTTCTGG ATTCAATCGAC TGTGGCCGGC TGGGTGTGGC GGACCGCTAT	1380
	CAGGACATAG CGTTGGCTAC CCGTGATATT GCTGAAGAGC TTGGCGGCGA ATGGGCTGAC	1440
10	CGCTTCCTCG TGCTTTACGG TATGCCGCT CCCGATTTCGC AGCGCATCGC CTTCTATCGC	1500
	CTTCTTGACG AGTTCTTCTG A	1521

## (2) INFORMATION FOR SEQ ID NO:32:

15           (i) SEQUENCE CHARACTERISTICS:  
               (A) LENGTH: 26 base pairs  
               (B) TYPE: nucleic acid  
               (C) STRANDEDNESS: single  
               (D) TOPOLOGY: linear

20           (ii) MOLECULE TYPE: DNA

25           (ix) FEATURE:  
               (A) NAME/KEY: -  
               (B) LOCATION: 1..26  
               (D) OTHER INFORMATION: /note= "oligonucleotide #18990"

30           (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:  
 GACCAGGGACA CGTATCCAGC CTCCGC

26

## (2) INFORMATION FOR SEQ ID NO:33:

35           (i) SEQUENCE CHARACTERISTICS:  
               (A) LENGTH: 28 base pairs  
               (B) TYPE: nucleic acid  
               (C) STRANDEDNESS: single  
               (D) TOPOLOGY: linear

40           (ii) MOLECULE TYPE: DNA

45           (ix) FEATURE:  
               (A) NAME/KEY: -  
               (B) LOCATION: 1..28  
               (D) OTHER INFORMATION: /note= "oligonucleotide #18991"

50           (xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:  
 GGAGGGCTGGA TACGTGTCCC GGTCTGCA

28

## (2) INFORMATION FOR SEQ ID NO:34:

55           (i) SEQUENCE CHARACTERISTICS:  
               (A) LENGTH: 7617 base pairs  
               (B) TYPE: nucleic acid  
               (C) STRANDEDNESS: single  
               (D) TOPOLOGY: linear

60           (ii) MOLECULE TYPE: DNA

65           (ix) FEATURE:

(A) NAME/KEY: -  
 (B) LOCATION: 1..7617  
 (D) OTHER INFORMATION: /note= "pGen-PGKgfo25R0"

5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:	
10	TCGAGGTCGA CGGTATCGAT TAGTCCAATT TGTAAAGAC AGGATATCAG TGGTCCAGGC	60
15	TCTAGTTTG ACTCAACAAT ATCACCAAGCT GAAGCCTATA GAGTACGAGC CATAGATAAA	120
20	ATAAAAGATT TTATTTAGTC TCCAGAAAAA GGGGGGAATG AAAGACCCC CCTGTAGGTT	180
25	TGGCAAGCTA GCTTAAGTAA CGCCATTITG CAAGGCATGG AAAAATACAT AACTGAGAAT	240
30	AGAGAAGTTC AGATCAAGGT CAGGAACAGA TGGAACAGCT GAATATGGC CAAACAGGAT	300
35	ATCTGTGGTA AGCAGTTCT GCCTCCGCTC AGGGCCAAGA ACAGATGGAA CAGCTGAATA	360
40	TGGGCCAAC AGGATATCTG TGGTAAGCAG TTCTGCCTCC GGCTCAGGGC CAAGAACAGA	420
45	TGGTCCCCAG ATGCGGTCCA GCCCTCAGCA GTTTCTAGAG AACCATCAGA TGTTTCCAGG	480
50	GTGCCCCAAG GACCTGAAAT GACCCTGTGC CTTATTTGAA CTAACCAATC AGTCGCTTC	540
55	TCGCTTCTGT TCGCGCGCTT CTGCTCCCCG AGCTCAATAA AAGAGCCAC AACCCCTCAC	600
60	TCGGGGCGCC AGTCCTCCGA TTGACTGAGT CGCCCGGGTA CCCGTGTATC CAATAAACCC	660
65	TCTTGCAGTT GCATCCGACT TGTGGTCTCG CTGTTCTTG GGAGGGTCTC CTCTGAGTGA	720
	TTGACTACCC GTCAGCGGGG GTCTTCATT TGGGGCTCG TCCGGGATCG GGAGACCCCT	780
	GCCCAGGGAC CACCGACCCA CCACCGGGAG GTAAGCTGGC CAGCAACTTA TCTGTGTCTG	840
	TCCGATTGTC TAGTGTCTAT GACTGATTT ATGCGCTGC GTCGGTACTA GTTAGCTAAC	900
	TAGCTCTGTA TCTGGCGGAC CCGTGGTGGA ACTGACGAGT TCGGAACACCC CGGCCGCAAC	960
	CCTGGGAGAC GTCCCAGGGA CTTCGGGGGC CGTTTTGTG GCCCGACCTG AGTCCAAAAAA	1020
	TCCCGATCGT TTTGGACTCT TTGGTGCACC CCCCTTAGAG GAGGGATATG TGGTTCTGGT	1080
	AGGAGACGAG AACCTAAAAC AGTTCCGCC TCCGTCTGAA TTTTGCTT CGGTTGGGA	1140
	CCGAAGCCGC GCCGCGCGTC TTGTCTGCTG CAGCATCGTT CTGTGTGTC TCTGTCTGAC	1200
	TGTGTTCTG TATTTGTCTG AGAATATGGG CCAGACTGTT ACCACTCCCT TAAGTTTGAC	1260
	CTTAGGTCACT TGGAAAGATG TCGAGCGGAT CGCTCACAAAC CAGTCGGTAG ATGTCAAGAA	1320
	GAGACGTTGG GTTACCTTCT GCTCTGCAGA ATGGCCAACC TTTAACGTG GATGGCCGCG	1380
	AGACGGCACC TTTAACCGAG ACCTCATCAC CCAGGTTAAG ATCAAGGTCT TTTCACCTGG	1440
	CCCGCATGGA CACCCAGACC AGGTCCCCTA CATCGTGACC TGGGAAGCCT TGGTTTTGA	1500
	CCCCCTCCCTGGTCAAGC CCTTTGTACA CCCTAACGCT CCGCCTCCCTC TTCTCCATC	1560
	CGCCCCGTCT CTCCCCCTTG AACCTCTCG TTCGACCCCG CCTCGATCCT CCCTTATCC	1620
	AGCCCTCACT CCTTCTCGAC GGTATACAGA CATGATAAGA TACATTGATG AGTTGGACA	1680
	AACCACAACT AGAATGCACT GAAAAAAATG CTTTATTTGT GAAATTTGTG ATGCTATTGC	1740
	TTTATTTGTA ACCATTATAA GCTGCAATAA ACAAGTTGGG GTGGGGCAAG AACTCCAGCA	1800
	TGAGATCCCC GCGCTGGAGG ATCATCCAGC CGCGAACGT GGCGAGAAAG GAAGGGAAGA	1860

	AAGCGAAAGG	AGCGGGCGCT	AGGGCGCTGG	CAAGTGTAGC	GGTCACGCTG	CGCGTAACCA	1920
	CCACACCCGC	CGCGCTTAAT	GCGCCGCTAC	AGGGCGCGTG	GGGATAACCCC	CTAGAGCCCC	1980
5	AGCTGGTTCT	TTCCGCCTCA	GAAGCCATAG	AGCCCACCGC	ATCCCCAGCA	TGCCCTGCTAT	2040
	TGTCTTCCA	ATCCTCCCCC	TTGCTGTCT	GCCCCACCCC	ACCCCCCAGA	ATAGAATGAC	2100
10	ACCTACTCAG	ACAATGCGAT	GCAATTTCCT	CATTTTATTA	GGAAAGGACA	GTGGGAGTGG	2160
	CACCTTCCAG	GGTCAAGGAA	GGCACGGGGG	AGGGGCAAAC	AACAGATGGC	TGGCAACTAG	2220
	AAGGCACAGT	CGAGGCTGAT	CAGCGAGCTC	TAGCATTAG	GTGACACTAT	AGAATAGGGC	2280
15	CCTCTAGATG	CATGCTCGAG	CGGCCGCCAG	TGTGATGGAT	ATCTGCAGAA	TTCTCAGAAG	2340
	AACTCGTCAA	GAAGGCGATA	GAAGGCGATG	CGCTGCGAAT	CGGGAGCGGC	GATACCGTAA	2400
20	AGCACGAGGA	AGCGGTCAAGC	CCATTGCCCG	CCAAGCTCTT	CAGCAATATC	ACGGGTAGCC	2460
	AACGCTATGT	CCTGATAGCG	GTCCGCCACA	CCCAGCCGGC	CACAGTCGAT	GAATCCAGAA	2520
	AAGCGGCCAT	TTTCCACCAT	GATATTGGC	AAGCAGGCAT	CGCCATGGGT	CACGACGAGA	2580
25	TCCTCGCCGT	CGGGCATGCG	CGCCTTGAGC	CTGGCGAACAA	GTTCGGCTGG	CGCGAGCCCC	2640
	TGATGCTCTT	CGTCCAGATC	ATCCTGATCG	ACAAGACCGG	CTTCCATCCG	AGTACGTGCT	2700
	CGCTCGATGC	GATGTTTCGC	TTGGTGGTCG	AATGGCAGG	TAGCCGGATC	AAGCGTATGC	2760
30	AGCCGCCGCA	TTGCATCAGC	CATGATGGAT	ACTTTCTCGG	CAGGAGCAAG	GTGAGATGAC	2820
	AGGAGATCCT	GCCCCGGCAC	TTCGCCCAAT	AGCAGCCAGT	CCCTTCCCAC	TTCAGTGACA	2880
35	ACGTCGAGCA	CAGCTGCGCA	AGGAACGCC	GTCGTGGCCA	GCCACGATAG	CCGCGCTGCC	2940
	TCGTCTGCA	TTTCATTCAAG	GGCACCCGAC	AGGTCGGTCT	TGACAAAAAG	AACCGGGCGC	3000
40	CCCTCGCTG	ACAGCCGGAA	CACGGCGCA	TCAGAGCAGC	CGATTGTCTG	TTGTGCCAG	3060
	TCATAGCCGA	ATAGCCTCTC	CACCCAAGCG	GCCGGAGAAC	CTGCGTGCAA	TCCATCTTGT	3120
	TCGATAGCAC	CAGCACCGTT	GTACAGTTCA	TCCATGCCAT	GTGTAATCCC	AGCAGCTGTT	3180
45	ACAAAATCAA	GAAGGACCAT	GTGGTCTCTC	TTTCGTTGG	GATCTTCGA	AAGGGCAGAT	3240
	TGTGTGGACA	GGTAATGGTT	GTCTGGAAA	AGGACAGGGC	CATCGCCAAT	TGGAGTATTT	3300
	TGTTGATAAT	GGTCTGCTAG	TTGAACCCTT	CCATCTCAA	TGTTGTGGCG	GGTCTTGAAG	3360
50	TTCACTTTGA	TTCCATTCTT	TTGTTGTCT	GCCATGATGT	ATACATTGTG	TGAGTTATAG	3420
	TTGTATTCCA	ATTGTGTCC	CAGAATGTTG	CCATCTTCCT	TGAAGTCAT	ACCTTTAAC	3480
55	TCGATTCTAT	TAACAAGGGT	ATCACCTCA	AACTGACTT	CAGCACGTGT	CTTGTAGTTG	3540
	CCGTCATCTT	TGAAGAAGAT	GGTCCTTCC	TGTACATAAC	CTTCGGGCAT	GGCACTCTTG	3600
60	AAAAAGTCAT	GCCGTTTCAT	ATGATCCGGG	TATCTTGAAA	AGCATTGAAC	ACCATAGCAC	3660
	AGAGTAGTGA	CTAGTGTGG	CCATGGAACA	GGCAGTTGC	CACTAGTGCA	GATGAACCTC	3720
	AGGGTAAGTT	TTCCGTATGT	TGCATCACCT	TCACCCCTCTC	CACTGACAGA	GAACCTGTGG	3780
65	CCGTTAACAT	CACCATCTAA	TTCAACAAGA	ATTGGGACAA	CTCCAGTGAA	GAGTTCTTCT	3840
	CCTTTGCTAG	CCATTCTTG	CGCGCCCGCG	GAGGCTGGAT	ACGTGTCCCG	GTCTGCAGGT	3900
	CGAAAGGCC	GGAGATGAGG	AAGAGGAGAA	CAGCGCGCA	GACGTGCGCT	TTTGAAGCGT	3960

	GCAGAATGCC	GGGCTCCGGA	GGACCTTCGC	GCCC GCCCG	CCCCTGAGCC	CGCCCCGTGAG	4020
	CCCGCCCCCG	GACCCACCCC	TTCCCAGCCT	CTGAGCCCAG	AAAGCGAAGG	AGCCAAGCTG	4080
5	CTATTGGCCG	CTGCCCCAAA	GGCCTACCCG	CTTCCATTGC	TCAGCGGTGC	TGTCCATCTG	4140
	CACGAGACTA	GTGAGACGTG	CTACTTCCAT	TTGTCACGTC	CTGCACGACG	CGAGCTGCGG	4200
10	GGCGGGGGGG	AACTTCCTGA	CTAGGGGAGG	AGTAGAAGGT	GGCGCGAAGG	GGCCACCAAA	4260
	GAAGGGAGCC	GGTTGGCGCT	ACCGGTGGAT	GTGGAATGTG	TGCGAGGCCA	GAGGCCACTT	4320
	GTGTAGCGCC	AACTGCCAGC	GGGGCTGCTA	AAGCGCATGC	TCCAGACTGC	CTTGGGAAAA	4380
15	GCGCCTCCCC	TACCCGGTAG	AATTGATAT	CAAGCTTATC	GATACCGTCG	AGATCTCCCG	4440
	ATCCGTCGAG	GTCGACGGTA	TCGATTAGTC	CAATTGTTA	AAGACAGGAT	ATCAGTGGTC	4500
20	CAGGCTCTAG	TTTGACTCA	ACAATATCAC	CAGCTGAAGC	CTATAGAGTA	CGAGCCATAG	4560
	ATAAAATAAA	AGATTTTATT	TAGTCTCCAG	AAAAAGGGGG	GAATGAAAGA	CCCCACCTGT	4620
	AGGTTTGGCA	AGCTAGCTTA	AGTAACGCCA	TTTGCAAGG	CATGGAAAAA	TACATAACTG	4680
25	AGAATAGAGA	AGTTCAGATC	GGGATCCCAA	TTCTTTCGGA	CTTTGAAAG	TGATGGTGGT	4740
	GGGGGAAGGA	TTCGAACCTT	CGAAGTCGAT	GACGGCAGAT	TTAGAGTCTG	CTCCCTTTGG	4800
30	CCGCTCGGGA	ACCCCACAC	GGGTAAATGCT	TTTACTGGCC	TGCTCCCTTA	TCGGGAAGCG	4860
	GGGGCGCATCA	TATCAAATGA	CGCGCCGCTG	TAAGTGTAA	CGTTGAGAAA	GAATTGGGAT	4920
	CCCGATCAAG	GTCAGGAACA	GATGGAACAG	CTAGAGAAC	ATCAGATGTT	TCCAGGGTGC	4980
35	CCCAAGGACC	TGAAATGACC	CTGTGCCCTA	TTTGAACCAA	CCAATCAGTT	CGCTTCTCGC	5040
	TTCTGTTCGC	GCGCTTCTGC	TCCCCGAGCT	CAATAAAAGA	GCCCACAACC	CCTCACTCGG	5100
40	GGGCCAGTC	CTCCGATTGA	CTGAGTCGCC	CGGGTACCCG	TGTATCCAAT	AAACCCCTTT	5160
	GCAGTTGCAT	CCGACTTGTG	GTCTCGCTGT	TCCTTGGGAG	GGTCTCCTCT	GAGTGATTGA	5220
	CTACCCGTCA	GGGGGGGTCT	TTCACCCAGA	TTTGGAACT	TACTGTCTTC	TTGGGACCTG	5280
45	CAGCCCCGGG	GATCCACTAG	TTCTAGAGCG	GCCGCCACCG	CGGTGGATT	TGCCTCGCGC	5340
	GTTTCGGTGA	TGACGGTGAA	AACCTCTGAC	ACATGCAGCT	CCCGGAGACG	GTCACAGCTT	5400
50	GTCTGTAAGC	GGATGCCGGG	AGCAGACAAG	CCCGTCAGGG	CGCGTCAGCG	GGTGTGGCG	5460
	GGTGTGGGGG	CGCAGCCATG	ACCCAGTCAC	GTAGCGATAG	CGGAGTGTAT	ACTGGCTTAA	5520
	CTATGCGGCA	TCAGAGCAGA	TTGTACTGAG	AGTGCACCAT	ATGCGGTGTG	AAATACCGCA	5580
55	CAGATGCGTA	AGGAGAAAAT	ACCGCATCG	GCGCTTCC	GCTTCTCGC	TCACTGACTC	5640
	GCTGCGCTCG	GTGCGTTGGC	TGCGCGAGC	GGTATCAGCT	CACTCAAAGG	CGGTAATACG	5700
60	GTTATCCACA	GAATCAGGGG	ATAACGCAGG	AAAGAACATG	TGAGCAAAG	GCCAGCAAA	5760
	GGCCAGGAAC	CGTAAAAAGG	CCGCGTTGCT	GGCGTTTTTC	CATAGGCTCC	GCCCCCTGA	5820
	CGAGCATCAC	AAAAATCGAC	GCTCAAGTCA	GAGGTGGCGA	AACCCGACAG	GACTATAAAG	5880
65	ATACCAAGCG	TTTCCCCCTG	GAAGCTCCCT	CGTGCCTCT	CCTGTTCCGA	CCCTGCCGCT	5940
	TACCGGATAC	CTGTCGGCCT	TTCTCCCTTC	GGGAAGCGTG	GCGCTTCTC	AATGCTCACG	6000
	CTGTAGGTAT	CTCAGTTCGG	TGTAGGTCGT	TCGCTCCAAG	CTGGGCTGTG	TGCACGAACC	6060

	CCCCGTTCA G	CCCGACCGCT GCGCCTTATC CGGTAACAT CGTCTTGAGT CCAACCCGGT	6120
	AAGACACGAC TTATGCCAC TGGCAGCAGC CACTGGTAAC AGGATTAGCA GAGCGAGGTA	6180	
5	TGTAGGGCGGT GCTACAGAGT TCTTGAAGTG GTGGCCTAAC TACGGCTACA CTAGAAGGAC	6240	
	AGTATTTGGT ATCTGCGCTC TGCTGAAGCC AGTTACCTTC GGAAAAAGAG TTGGTAGCTC	6300	
10	TTGATCCGGC AAACAAACCA CCGCTGGTAG CGGTGGTTTT TTTGTTTGCA AGCAGCAGAT	6360	
	TACGCGCAGA AAAAAAGGAT CTCAAGAAGA TCCTTTGATC TTTTCTACGG GGTCTGACGC	6420	
	TCAGTGGAAC GAAAACTCAC GTTAAGGGAT TTTGGTCATG AGATTATCAA AAAGGATCTT	6480	
15	CACCTAGATC CTTTAAATT AAAAATGAAG TTTTAAATCA ATCTAAAGTA TATATGAGTA	6540	
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30	TGGTATGGCT TCATTCAGCT CCGGTTCCA ACGATCAAGG CGAGTTACAT GATCCCCAT	6960	
	GTTGTGCAAA AAAGCGGTTA GCTCCTTCGG TCCTCCGATC GTTGTAGAA GTAAGTTGGC	7020	
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	TTTACTTTT ACCAGCGTTT CTGGGTGAGC AAAAAACAGGA AGGAAAATG CCGCAAAAAA	7380	
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50	TAAACAAATA GGGGTCCGC GCACATTTC CCGAAAATG CCACCTGACG TCTAAGAAC	7560	
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## (2) INFORMATION FOR SEQ ID NO:35:

(i) SEQUENCE CHARACTERISTICS:  
 5 (A) LENGTH: 15581 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: DNA

15 (ix) FEATURE:  
 (A) NAME/KEY: -  
 (B) LOCATION: 1..15581  
 (D) OTHER INFORMATION: /note= "pNLnSG11"

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

20	TGGAAGGGCT AATTTGGTCC CAAAAAAAGAC AAGAGATCCT TGATCTGTGG ATCTACCACA	60
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25	TGACCTTTGG ATGGTGCTTC AAGTTAGTAC CAGTTGAACC AGAGCAAGTA GAAGAGGCCA	180
	AATAAGGAGA GAAGAACAGC TTGTTACACC CTATGAGCCA GCATGGGATG GAGGACCCGG	240
	AGGGAGAAGT ATTAGTGTGG AAGTTTGACA GCCTCCTAGC ATTCGTCAC ATGGCCCGAG	300
30	AGCTGCATCC GGAGTACTAC AAAGACTGCT GACATCGAGC TTTCTACAAG GGACTTTCCG	360
	CTGGGGACTT TCCAGGGAGG TGTGGCCTGG GCGGGACTGG GGAGTGGCGA GCCCTCAGAT	420
35	GCTACATATA AGCAGCTGCT TTTTGCCCTGT ACTGGGTCTC TCTGGTTAGA CCAGATCTGA	480
	GCCTGGGAGC TCTCTGGCTA ACTAGGGAAC CCACTGCTTA AGCCTCAATA AAGCTTGCCT	540
	TGAGTGCTCA AAGTAGTGTG TGCCCGTCTG TTGTGTGACT CTGGTAACTA GAGATCCCTC	600
40	AGACCCCTTT AGTCAGTGTG GAAAATCTCT AGCAGTGGCG CCCGAACAGG GACTTGAAAG	660
	CGAAAGTAAA GCCAGAGGGAG ATCTCTCGAC GCAGGACTCG GCTTGCTGAA GCGCGCACGG	720
45	CAAGAGGCCA GGGGCGGCAGA CTGGTGAGTA CGCCAAAAAT TTTGACTAGC GGAGGCTAGA	780
	AGGAGAGAGA TGGGTGCGAG AGCGTCGGTA TTAAGCGGGG GAGAATTAGA TAAATGGAA	840
	AAAATTCCGGT TAAGGCCAGG GGGAAAGAAA CAATATAAAC TAAAACATAT AGTATGGCA	900
50	AGCAGGGAGC TAGAACGATT CGCAGTTAAT CCTGGCCTTT TAGAGACATC AGAAGGCTGT	960
	AGACAAATAC TGGGACAGCT ACAACCATCC CTTCAGACAG GATCAGAAGA ACTTAGATCA	1020
55	TTATATAATA CAATAGCAGT CCTCTATTGT GTGCATCAAA GGATAGATGT AAAAGACACC	1080
	AAGGAAGCCT TAGATAAGAT AGAGGAAGAG CAAAACAAAA GTAAGAAAAA GGCACAGCAA	1140
	GCAGCAGCTG ACACAGGAAA CAACAGCCAG GTCAGCCAAA ATTACCCAT AGTGCAGAAC	1200
60	CTCCAGGGGC AAATGGTACA TCAGGCCATA TCACCTAGAA CTTTAAATGC ATGGGTAAAA	1260
	GTAGTAGAAG AGAAGGCTTT CAGCCCAGAA GTAATACCCA TGTTTTCAGC ATTATCAGAA	1320
65	GGAGGCCACCC CACAAGATTT AAATACCATG CTAAACACAG TGGGGGGACA TCAAGCAGCC	1380
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	CCACCAGAAG AGAGCTTCAG GTTGGGGAA GAGACAACAA CTCCTCTCA GAAGCAGGAG	2220
25	CCGATAGACA AGGAACGTGA TCCTTAGCT TCCCTCAGAT CACTCTTGG CAGCGACCCC	2280
	TCGTCACAAT AAAGATAGGG GGGCAATTAA AGGAAGCTCT ATTAGATACA GGAGCAGATG	2340
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	AAGCTATAGG TACAGTATTAA GTAGGACCTA CACCTGTCAA CATAATTGGA AGAAATCTGT	2520
35	TGACTCAGAT TGGCTGCACT TTAAATTTTC CCATTAGTCC TATTGAGACT GTACCAGTAA	2580
	AATTAAAGCC AGGAATGGAT GGCCCAAAAG TTAAACAATG GCCATTGACA GAAGAAAAAA	2640
	TAAAAGCATT AGTAGAAATT TGTACAGAAA TGGAAAAGGA AGGAAAAATT TCAAAAATTG	2700
40	GGCCTGAAAA TCCATACAAT ACTCCAGTAT TTGCCATAAA GAAAAAAGAC AGTACTAAAT	2760
	GGAGAAAATT AGTAGATTTC AGAGAACTTA ATAAGAGAAC TCAAGATTTC TGGGAAGTTC	2820
45	AATTAGGAAT ACCACATCCT GCAGGGTTAA AACAGAAAAA ATCAGTAACA GTACTGGATG	2880
	TGGCGATGC ATATTTTCA GTTCCCTTAG ATAAAGACTT CAGGAAGTAT ACTGCATTAA	2940
	CCATACCTAG TATAAACAAAT GAGACACCAG GGATTAGATA TCAGTACAAT GTGCTTCCAC	3000
50	AGGGATGGAA AGGATCACCA GCAATATTCC AGTGTAGCAT GACAAAAATC TTAGAGCCTT	3060
	TTAGAAAACA AAATCCAGAC ATAGTCATCT ATCAATACAT GGATGATTG TATGTAGGAT	3120
55	CTGACTTAGA AATAGGGCAG CATAGAACAA AAATAGAGGA ACTGAGACAA CATCTGTTGA	3180
	GGTGGGGATT TACCACACCA GACAAAAAAC ATCAGAAAGA ACCTCCATTC CTTTGGATGG	3240
	GTTATGAACT CCATCCTGAT AAATGGACAG TACAGCCTAT AGTGCTGCCA GAAAAGGACA	3300
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	ACACAACAAA TCAGAAGACT GAGTTACAAG CAATTCACT AGCTTGTGAG GATTGGGAT	4020
15	TAGAAGTAAA CATAGTGACA GACTCACAAAT ATGCATTGGG AATCATTCAA GCACAACCAG	4080
	ATAAAGAGTGA ATCAGAGTTA GTCAGTCAA TAATAGAGCA GTTAATAAAA AAGGAAAAG	4140
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35	TAAGGTGCAG AAAGAATATG CATTCTTTA TAAACTTGAT ATAGTACCAA TAGATAATAC	6780
	CAGCTATAGG TTGATAAGTT GTAACACCTC AGTCATTACA CAGGCCTGTC CAAAGGTATC	6840
40	CTTTGAGCCA ATTCCCATAAC ATTATTGTGC CCCGGCTGGT TTTGCGATTTC TAAAATGTAA	6900
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5	AGTGTAGAG TGGAGGTTTG ACAGCCGCCT AGCATTTCAT CACGTGGCCC GAGAGCTGCA	10080
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25	AGTTTGGAG TTCCAGACCA GCCTGACCAA CATGGAGAAA CCCCTTCTCT GTGTATTTT	10680
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30	TTCCCTACT CTGATACAC AAGAACATC AGCACAGAGG AAGACTTCG TGATCAAATG	10800
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35	GGGCTCAGTC CCCAAGACAT AAACACCCAA GACATAAACCA CCCAACAGGT CCACCCGCC	10980
	TGCTGCCAG GCAGAGCCGA TTCACCAAGA CGGGAATTAG GATAGAGAAA GAGTAAGTCA	11040
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	TGAAAGCGTA GGGAGTCGAA GGTGTCTTT TGCGCCGAGT CAGTTCTGG GTGGGGGCCA	11220
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5	ACGGTTATCC ACAGAACAGC GGGATAACGC AGGAAAGAAC ATGTGAGCAA AAGGCCAGCA	12180
	AAAGGCCAGG AACCGTAAAA AGGCCGCGTT GCTGGCGTTT TTCCATAGGC TCCGCCCCC	12240
10	TGACGAGCAT CACAAAATC GACGCTCAAG TCAGAGGTGG CGAAACCCGA CAGGACTATA	12300
	AAGATACCAG GCGTTTCCCC CTGGAAGCTC CCTCGTGCAG TCTCCTGTT CGACCCCTGCC	12360
	GCTTACCGGA TACCTGTCCG CCTTCTCCC TTCGGGAAGC GTGGCGCTTT CTCAATGCTC	12420
15	ACGCTGTAGG TATCTCAGTT CGGTGTAGGT CGTCGCTCC AAGCTGGGCT GTGTGCACGA	12480
	ACCCCCCGTT CAGCCCGACC GCTGCGCCTT ATCCGGTAAC TATCGTCTTG AGTCCAACCC	12540
20	GGTAAGACAC GACTTATCGC CACTGGCAGC AGCCACTGGT AACAGGATTAA GCAGAGCGAG	12600
	GTATGTAGGC GGTGCTACAG AGTTCTGAA GTGGTGGCCT AACTACGGCT ACACTAGAAG	12660
	GACAGTATTT GGTATCTGCG CTCTGCTGAA GCCAGTTACC TTCGGAAAAA GAGTTGGTAG	12720
25	CTCTTGATCC GGCAAAACAAA CCACCGCTGG TAGCGGTGGT TTTTTTGTGTT GCAAGCAGCA	12780
	GATTACGCGC AGAAAAAAAG GATCTCAAGA AGATCCTTTG ATCTTTCTA CGGGGTCTGA	12840
30	CGCTCAGTGG AACGAAAAGT CACGTTAAGG GATTTGGTC ATGAGATTAT CAAAAAGGAT	12900
	CTTCACCTAG ATCCTTTAA ATTAAAAATG AAGTTTTAAA TCAATCTAAA GTATATATGA	12960
	GTAAACATTGG TCTGACAGTT ACCAATGCTT AATCAGTGAG GCACCTATCT CAGCGATCTG	13020
35	TCTATTCGT TCATCCATAG TTGCCTGACT CCCCGTCGTG TAGATAACTA CGATACGGGA	13080
	GGGTTTACCA TCTGGCCCCA GTGCTGCAAT GATACCGCGA GACCCACGCT CACCGCTCC	13140
40	AGATTTATCA GCAATAAAACC AGCCAGCCGG AAGGGCCGAG CGCAGAAGTG GTCCTGCAAC	13200
	TTTATCCGCC TCCATCCAGT CTATTAATTG TTGCCGGGAA GCTAGAGTAA GTAGTTGCC	13260
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45	GTGGGTATG GCTTCATTCA GCTCCGGTTC CCAACGATCA AGGCGAGTTA CATGATCCCC	13380
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50	GGCCGCAGTG TTATCACTCA TGGTTATGGC AGCACTGCAT AATTCTCTTA CTGTCATGCC	13500
	ATCCGTAAGA TGCTTTCTG TGACTGGTGA GTACTCAACC AAGTCATTCT GAGAATAGTG	13560
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60	ATCTTTTACT TTCACCAGCG TTTCTGGGTG AGCAAAAACA GGAAGGCAAATGCCGCAA	13800
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65	AAATAAACAA ATAGGGGTTT CGCGCACATT TCCCCGAAAA GTGCCACCTG ACGTCTAAGA	13980
	AACCATTATT ATCATGACAT TAACCTATAA AAATAGGCGT ATCACGAGGC CCTTCGTCT	14040
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	CGTCAGCGGG TGTTGGCGGG TGTGCGGGCG CAGCCATGAC CCAGTCACGT AGCGATAGCG	14220
5	GAGTGTACTG GCTTAACTAT CGGGCATCAG AGCAGATTGT ACTGAGAGTG CACCATATGC	14280
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10	TCAGGCTGCG CAACTGTTGG GAAGGGCGAT CGGTGCGGGC CTCTTCGCTA TTACGCCAGC	14400
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20	TGTATACAAA ATCTAGGCCA GTCCAGCAGA GCCTAAAGGT AAAAAATAAA ATAATAAAAA	14700
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45	CTCCACGTGC ACATGGCCGG AGGAACCTGCC ATGTCGGAGG TGCAAGCACA CCTGCGCATC	15480
	AGAGTCCTTG GTGTGGAGGG AGGGACCAGC GCAGCTTCCA GCCATCCACC TGATGAACAG	15540
50	AACCTAGGGA AAGCCCCAGT TCTACTTACA CCAGGAAAGG C	15581

## (2) INFORMATION FOR SEQ ID NO:36:

5 (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 74 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

10 (ii) MOLECULE TYPE: DNA

15 (ix) FEATURE:  
(A) NAME/KEY: -  
(B) LOCATION: 1..74  
(D) OTHER INFORMATION: /note= "primer #17982"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

20 GGGCGTACG GAGCGCTCCG AATTCCGTAC CGTTAACG GGCCCTCTCG AGTCCGTTGT 60  
ACAGTTCATC CATG 74

25 (2) INFORMATION FOR SEQ ID NO:37:

30 (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 66 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

35 (ii) MOLECULE TYPE: DNA

40 (ix) FEATURE:  
(A) NAME/KEY: -  
(B) LOCATION: 1..66  
(D) OTHER INFORMATION: /note= "primer #17983"

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

45 GGGGAATTG GCGCGCGTAC GAAAGCGCTA GCTGAGCAAG AAATGGCTAG CAAAGGAGAA 60  
GAACTC 66

WHAT IS CLAIMED IS:

- 1           1. An isolated nucleic acid that encodes an  
2         engineered *Aequorea victoria* fluorescent protein, wherein the  
3         protein encoded by the isolated nucleic acid is selected from  
4         the group that consists of:
  - 5           a. a protein that has leucine at amino acid position  
6           65, and wherein said protein has a cellular  
7           fluorescence that is at least five times greater  
8           than the cellular fluorescence of wild type *Aequorea*  
9           *victoria* green fluorescent protein;
  - 10          b. a protein that has leucine at amino acid position 65  
11         and threonine at position 168, and wherein said  
12         protein has a cellular fluorescence that is at least  
13         five times greater than wild type *Aequorea victoria*  
14         green fluorescent protein;
  - 15          c. a protein that has leucine at amino acid position 65  
16         threonine at position 168, and cysteine at position  
17         66, wherein said protein has a cellular fluorescence  
18         that is at least five times greater than the  
19         cellular fluorescence of wild type *Aequorea victoria*  
20         green fluorescent protein;
  - 21          d. A blue fluorescent protein that has histidine at  
22         amino acid position 67, leucine at position 65 and  
23         has a cellular fluorescence that is at least five  
24         times greater than that of BFP(Tyr<sub>67</sub>→His);
  - 25          e. a blue fluorescent protein that has histidine at  
26         amino acid position 67, alanine at amino acid  
27         position 164 and has a cellular fluorescence that is  
28         at least five times greater than that of  
29         BFP(Tyr<sub>67</sub>→His);
  - 30          f. a blue fluorescent protein that has histidine at  
31         amino acid position 67, leucine at amino acid  
32         position 65, alanine at amino acid position 164 and  
33         has a cellular fluorescence that is at least five  
34         times greater than that of BFP(Tyr<sub>67</sub>→His).

1           2. An isolated nucleic acid of claim 1, which  
2 encodes an engineered *Aequorea victoria* green fluorescent  
3 protein ("GFP") having a cellular fluorescence that is at  
4 least five times greater than that of wild type GFP, wherein  
5 the engineered GFP has a leucine at amino acid position 65.

1           3. An isolated nucleic acid according to claim 2,  
2 wherein the nucleic acid further encodes a threonine at amino  
3 acid position 168.

1           4. An isolated nucleic acid according to claim 3,  
2 wherein the nucleic acid further encodes a cysteine at amino  
3 acid position 66.

1           5. An isolated nucleic acid of claim 1 that  
2 encodes an engineered blue fluorescent protein ("BFP") that  
3 has histidine at amino acid position 67 and leucine at  
4 position 65, and has a cellular fluorescence that is at least  
5 five times greater than that of BFP( $Tyr_{67} \rightarrow His$ ).

1           6. An isolated nucleic acid of claim 1 that  
2 encodes an engineered blue fluorescent protein ("BFP") that  
3 has histidine at amino acid position 67 and alanine at amino  
4 acid position 164, and has a cellular fluorescence that is at  
5 least five times greater than that of BFP( $Tyr_{67} \rightarrow His$ ).

1           7. An isolated nucleic acid according to claim 6,  
2 wherein the nucleic acid further encodes leucine at amino acid  
3 position 65.

1           8. A transformed cell that expresses a protein  
2 encoded by a nucleic acid of claim 1.

1           9. A vector comprising a nucleic acid of claim 1.

1           10. A transformed cell comprising a vector of  
2 claim 9.

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1           11. A transformed cell that expresses a protein  
2 encoded by the nucleic acid of claim 1 fused to a protein  
3 encoded by a second nucleic acid of interest.

1           12. An isolated engineered *Aequorea victoria* green  
2 fluorescent protein ("GFP") wherein the engineered GFP  
3 comprises leucine at amino acid position 65, said engineered  
4 GFP having a cellular fluorescence that is at least five times  
5 greater than wild type GFP.

1           13. An isolated engineered *Aequorea victoria* green  
2 fluorescent protein ("GFP") according to claim 12, wherein the  
3 engineered GFP has threonine at amino acid position 168.

1           14. An isolated engineered *Aequorea victoria* green  
2 fluorescent protein ("GFP") according to claim 13, wherein the  
3 engineered GFP has cysteine at amino acid position 66.

1           15. An isolated blue fluorescent protein ("BFP")  
2 that comprises histidine at amino acid position 67 and leucine  
3 at amino acid position 65 and has a cellular fluorescence that  
4 is at least five times greater than that of BFP(Tyr<sub>67</sub>→His).

1           16. An isolated blue fluorescent protein ("BFP")  
2 that has a histidine at amino acid position 67 and an alanine  
3 at amino acid position 164, that has a cellular fluorescence  
4 that is at least five times greater than that of  
5 BFP(Tyr<sub>67</sub>→His).

1           17. An isolated blue fluorescent protein ("BFP")  
2 according to claim 16, wherein the BFP further has leucine at  
3 amino acid position 65.

1           18. A method of detecting and optionally isolating  
2 an engineered cell that contains a selected nucleic acid which  
3 encodes a selected protein or nucleic acid, comprising:  
4           a) stably introducing into a host cell in a population of  
5 host cells a vector that contains a first nucleic acid which

6       encodes a polypeptide selected from the group consisting of  
7       SG11, SG12, SG25, SB42, SB49, SB50 and a second nucleic acid  
8       which encodes a selected protein or nucleic acid, and  
9            b) detecting cells in the population of host cells that  
10      express SG11, SG12, SG25, SB42, SB49, or SB50, and  
11            c) optionally sorting cells that express SG11, SG12,  
12      SG25, SB42, SB49, or SB50 with a fluorescence-activated cell  
13      sorter to isolate individual cells that express said  
14      fluorescent protein.

1           19. A nucleic acid construct wherein a coding  
2       sequence selected from the group consisting of sequences that  
3       encode SG11, SG12, SG25, SB42, SB49, and SB50 is operably  
4       linked to a regulatory sequence of a selected gene.

1           20. A nucleic acid construct wherein a first coding  
2       sequence that encodes a selected polypeptide is fused using  
3       genetic engineering to a second coding sequence selected from  
4       the group consisting of sequences that encode SG11, SG12,  
5       SG25, SB42, SB49, and SB50, such that expression of the fused  
6       sequence yields a fluorescent hybrid protein in which the  
7       polypeptide encoded by the first coding sequence is fused to  
8       the polypeptide encoded by the second coding sequence.

1           21. A method of detecting and characterizing  
2       regulatory and coding sequence elements that regulate  
3       subcellular expression and targeting of proteins, comprising:  
4            a) expressing in an engineered cell, in the presence and  
5       absence of selected culture conditions and components, a  
6       nucleic acid wherein a first nucleic acid selected from the  
7       group consisting of nucleic acids that encode SG11, SG12,  
8       SG25, SB42, SB49, and SB50 is operably linked to a second  
9       nucleic acid derived from a selected gene;  
10           b) detecting the presence and subcellular localization of  
11       fluorescent signal.

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 97/07625

A. CLASSIFICATION OF SUBJECT MATTER					
IPC 6	C12N15/12	C12N15/62	C12N15/85	C12N15/86	C12N5/10
	C07K14/435	C12Q1/02	C12Q1/68	G01N33/50	G01N33/52

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 C12N C07K C12Q G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	<p>WO 97 11094 A (NOVONORDISK AS ;THASTRUP OLE (DK); TULLIN SOEREN (DK); POULSEN LAR) 27 March 1997</p> <p>see the whole document</p> <p>---</p>	1,2,5, 8-10,12, 15,18, 19,21
A	<p>PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, vol. 91, 1 December 1994, pages 12501-12504, XP000574454</p> <p>HEIM R ET AL: "WAVELENGTH MUTATIONS AND POSTTRANSLATIONAL AUTOXIDATION OF GREEN FLUORESCENT PROTEIN" cited in the application see page 12504, left-hand column, line 8 - right-hand column, line 8; table 1</p> <p>---</p> <p>-/-</p>	1-21

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

\* Special categories of cited documents :

- 'A' document defining the general state of the art which is not considered to be of particular relevance
- 'E' earlier document but published on or after the international filing date
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- 'O' document referring to an oral disclosure, use, exhibition or other means
- 'P' document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search  14 August 1997	Date of mailing of the international search report  26.08.97
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Hornig, H

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/07625

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	
A	NATURE, vol. 373, 23 February 1995, page 663/664 XP002029558 HEIM R ET AL: "IMPROVED GREEN FLUORESCENCE" cited in the application see the whole document ---	1-21	
A	BIO/TECHNOLOGY, vol. 13, no. 2, February 1995, pages 151-154, XP002033686 DELAGRAVE S ET AL: "RED-SHIFTED EXCITATION MUTANTS OF THE GREEN FLUORESCENT PROTEIN" cited in the application see table 1 ---	1-21	
A	FEBS LETTERS, vol. 367, 1 January 1995, pages 163-166, XP000579119 EHRIG T ET AL: "GREEN-FLUORESCENT PROTEIN MUTANTS WITH ALTERED FLUORENCE EXCITATION SPECTRA" see page 165, left-hand column, line 1 - right-hand column, line 37 ---	1-21	
A	TIBS TRENDS IN BIOCHEMICAL SCIENCES, vol. 20, November 1995, pages 448-455, XP000606919 CUBITT A B ET AL: "UNDERSTANDING, IMPROVING AND USING GREEN FLUORESCENT PROTEINS" see the whole document ---	1-21	
A	WO 95 07463 A (UNIV COLUMBIA ;WOODS HOLE OCEANOGRAPHIC INST (US); CHALFIE MARTIN) 16 March 1995 see the whole document ---	18-21	
P,X	GENE, vol. 173, no. 1, 1 July 1996, ELSEVIER SCIENCE PUBLISHERS,B.V.,AMSTERDAM,NL;, pages 33-38, XP002037756 B.P. CORMACK ET AL.: "FACS-optimized mutants of the green fluorescent protein (GFP)" see page 37, left-hand column, line 12 - right-hand column, line 4 ---	1,2, 8-10,12, 19	
1	P,A	WO 96 23810 A (UNIV CALIFORNIA ;TSIEN ROGER Y (US); HEIM ROGER (US)) 8 August 1996 see the whole document -----	1-21

**INTERNATIONAL SEARCH REPORT**

International Application No

PCT/US 97/07625

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WO 9711094 A	27-03-97	AU 4482996 A		09-04-97
WO 9507463 A	16-03-95	US 5491084 A AU 7795794 A CA 2169298 A EP 0759170 A		13-02-96 27-03-95 16-03-95 26-02-97
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